

GenCore version 5.1.6  
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OM nucleic - nucleic search, using sw model

Run on: August 14, 2003, 21:44:31 ; Search time 29.025 Seconds  
(without alignments)  
273.726 Million cell updates/sec

Title: US-10-074-620-1

Sequence: 1 ggcctggctgcacctgtta 18

Scoring table: OLIGO\_NUC  
Gapop 60.0 , Gapext 60.0

Searched: 569978 seqs, 220691566 residues

Word size : 0

Total number of hits satisfying chosen parameters: 1139956

Minimum DB seq length: 0  
Maximum DB seq length: 200000000

Post-processing: Listing first 120 summaries

Database :

Issued Patents\_NA:\*  
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3: /cgm2\_6/prodata/2/ina/6A\_COMB.seq:\*  
4: /cgm2\_6/prodata/2/ina/6B\_COMB.seq:\*  
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6: /cgm2\_6/prodata/2/ina/backfile1.seq:\*

Pred. No. is the number of results predicted by chance to have a  
score greater than or equal to the score of the result being printed,  
and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	16	88.9	20	1	US-08-197-791-28
2	16	88.9	50	1	US-08-171-389-546
3	16	88.9	50	1	US-08-123-936-546
4	16	88.9	50	2	US-08-475-228A-546
5	16	88.9	50	3	US-08-482-080A-546
6	16	88.9	50	4	US-09-354-947-546
7	16	88.9	50	5	PCT-US93-12388-546
8	16	88.9	50	5	PCT-US93-12388-546
9	16	88.9	50	5	PCT-US93-12388-546
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77	12	66.7	4015	3	US-08-974-549A-1	Sequence 1, Appli
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81	12	66.7	4015	4	US-09-572-423B-3	Sequence 1, Appli
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 115 12 66.7 31960 4 US-09-679-409-1  
 116 12 66.7 44037 3 US-09-103-840A-2  
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 118 11 61.1 20 3 US-08-450-905B-125  
 119 11 61.1 20 3 US-07-982-759F-125  
 120 11 61.1 20 4 US-09-422-978-5580

ALIGNMENTS

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 Sequence 4, Appl  
 Sequence 4, Appl  
 Sequence 5, Appl  
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 Sequence 3, Appl  
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 Sequence 1, Appl  
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 Sequence 125, App  
 Sequence 5580, Ap

RESULT 1

US-08-197-791-28

Sequence 26, Application US/08/197-791-28

Patent No. 5,567,772

GENERAL INFORMATION:

APPLICANT: Sierge, Joseph A.

APPLICANT: Mullinax, Rebecca L.

TITLE OF INVENTION: NOVEL POLYMERASE COMPOSITIONS AND USES

TITLE OF INVENTION: THEREOF

NUMBER OF SEQUENCES: 44

CORRESPONDENCE ADDRESSES:

ADDRESSEE: Limbach and Limbach

STREET: 2001 Ferry Building

CITY: San Francisco

STATE: CA

COUNTRY: USA

ZIP: 94111

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk

COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: Patent in Release #1.0, Version #1.25

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/197,791

FILING DATE:

CLASSIFICATION: 424

PRIOR APPLICATION DATA:

APPLICATION NUMBER: US 08/164,290

FILING DATE: 08-DEC-1993

ATTORNEY/AGENT INFORMATION:

NAME: Bortner, Scott R.

REGISTRATION NUMBER: 34,298

REFERENCE/DOCKET NUMBER: 20270 USA

TELEPHONE: 415-433-8216

TELEFAX: 415-433-4150

INFORMATION FOR SEQ ID NO: 28:

SEQUENCE CHARACTERISTICS:

LENGTH: 20 base pairs

TYPE: nucleic acid

STRANDEDNESS: single

TOPOLOGY: linear

MOLECULE TYPE: DNA (genomic)

HYPOTHETICAL: NO

ANTI-SENSE: NO

US-08-197-791-28

Query Match

88.9%; Score 16; DB 1; Length 20;

Best Local Similarity 100.0%; Pred. No. 1.2;  
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 GGGTGGTGCACCTGT 16  
 Db 1 GGGTGGTGCACCTGT 16

RESULT 2

US-08-171-389-546/C

Sequence 546, Application US/08/171389

Patent No. 5,567,772

GENERAL INFORMATION:

APPLICANT: Edwards, Cynthia A.

APPLICANT: Cantor, Charles R.

APPLICANT: Andrews, Beth M.

APPLICANT: Turin, Lisa M.

TITLE OF INVENTION: Sequence-Directed DNA Binding

TITLE OF INVENTION: Molecules, Compositions and Methods

NUMBER OF SEQUENCES: 641

CORRESPONDENCE ADDRESSES:

ADDRESSEE: Genelabs Technologies, Inc.

STREET: 505 Penobscot Drive

CITY: Redwood City

STATE: CA

COUNTRY: USA

ZIP: 94063

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk

COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: Patent in Release #1.0, Version #1.25

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/171,389

FILING DATE:

CLASSIFICATION: 435

PRIOR APPLICATION DATA:

APPLICATION NUMBER: US 08/123,936

FILING DATE: 17-SEP-1993

PRIOR APPLICATION DATA:

APPLICATION NUMBER: US 07/996,783

FILING DATE: 23-DEC-1992

PRIOR APPLICATION DATA:

APPLICATION NUMBER: US 07/723,618

FILING DATE: 27-JUN-1991

PRIOR APPLICATION DATA:

APPLICATION NUMBER: US 08/081,070

FILING DATE: 22-JUN-1993

ATTORNEY/AGENT INFORMATION:

NAME: Fabian, Gary R.

REGISTRATION NUMBER: 33,875

REFERENCE/DOCKET NUMBER: 4600-0175/G19P3

TELEPHONE: (415) 324-0880

TELEFAX: (415) 324-0960

INFORMATION FOR SEQ ID NO: 546:

SEQUENCE CHARACTERISTICS:

LENGTH: 50 base pairs

TYPE: nucleic acid

STRANDEDNESS: double

TOPOLOGY: linear

MOLECULE TYPE: DNA (genomic)

HYPOTHETICAL: NO

ORIGINAL SOURCE:

INDIVIDUAL ISOLATE: Epstein Barr virus L2 (start site)

INDIVIDUAL ISOLATE: 900211

US-08-171-389-546

Query Match

88.9%; Score 16; DB 1; Length 50;

Best Local Similarity 100.0%; Pred. No. 1.2;

Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 GGCTGGTGCACCTGT 16  
DB 41 GGCTGGTGCACCTGT 26

## RESULT 3

US-08-123-936-546/c  
Sequence 546, Application US/08123936  
Patent No. 5726014  
GENERAL INFORMATION:  
APPLICANT: Edwards, Cynthia A.  
APPLICANT: Cantor, Charles R.  
APPLICANT: Andrews, Beth M.  
APPLICANT: Turin, Lisa M.  
TITLE OF INVENTION: Screening Assay for the Detection of  
NUMBER OF SEQUENCES: 640  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Genelabs Technologies, Inc.  
STREET: 505 Penobscot Drive  
CITY: Redwood City  
STATE: CA  
COUNTRY: USA  
ZIP: 94063  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: Patentin Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/123,936  
FILING DATE:  
CLASSIFICATION: 435  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/996,783  
FILING DATE: 23-DEC-1992  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/723,618  
FILING DATE: 27-JUN-1991  
ATTORNEY/AGENT INFORMATION:  
NAME: Fabian, Gary R.  
REGISTRATION NUMBER: 33,875  
REFERENCE/DOCKET NUMBER: 4600-0075.32/G19P2  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: (415) 324-0880  
TELEFAX: (415) 324-0960  
INFORMATION FOR SEQ ID NO: 546:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 50 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: double  
TOPOLOGY: linear  
MOLECULE TYPE: DNA (genomic)  
HYPOTHETICAL: NO  
ORIGINAL SOURCE:  
INDIVIDUAL ISOLATE: Epstein Barr virus L2 (start site)  
INDIVIDUAL ISOLATE: 90021  
US-08-123-936-546

Query Match 88.9%; Score 16; DB 1; Length 50;  
Best Local Similarity 100.0%; Pred. No. 1.2;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 GGCTGGTGCACCTGT 16  
DB 41 GGCTGGTGCACCTGT 26

RESULT 4  
US-08-475-228A-546/c  
Sequence 546, Application US/08475228A  
Patent No. 5869241  
GENERAL INFORMATION:

APPLICANT: Edwards, Cynthia A.  
APPLICANT: Cantor, Charles R.  
APPLICANT: Andrews, Beth M.  
APPLICANT: Turin, Lisa M.  
APPLICANT: Fry, Kirk E.  
TITLE OF INVENTION: Sequence-Directed DNA Binding  
NUMBER OF SEQUENCES: 664  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Genelabs Technologies, Inc.  
STREET: 505 Penobscot Drive  
CITY: Redwood City  
STATE: CA  
COUNTRY: USA  
ZIP: 94063  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: Patentin Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/475,228A  
FILING DATE: 06-JUN-1995  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/123,936  
FILING DATE: 17-SEP-1993  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/996,783  
FILING DATE: 23-DEC-1992  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/723,618  
FILING DATE: 27-JUN-1991  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/081,070  
FILING DATE: 22-JUN-1993  
ATTORNEY/AGENT INFORMATION:  
NAME: Stratford, Carol A.  
REGISTRATION NUMBER: 34,444  
REFERENCE/DOCKET NUMBER: 4600-0175.21/G19P3D2  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: (415) 324-0880  
TELEFAX: (415) 324-0960  
INFORMATION FOR SEQ ID NO: 546:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 50 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: double  
TOPOLOGY: linear  
MOLECULE TYPE: DNA (genomic)  
HYPOTHETICAL: NO  
ORIGINAL SOURCE:  
INDIVIDUAL ISOLATE: Epstein Barr virus L2 (start site)  
INDIVIDUAL ISOLATE: 90021  
US-08-475-228A-546

Query Match 88.9%; Score 16; DB 2; Length 50;  
Best Local Similarity 100.0%; Pred. No. 1.2;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 GGCTGGTGCACCTGT 16  
DB 41 GGCTGGTGCACCTGT 26

RESULT 5  
US-08-482-080A-546/c  
Sequence 546, Application US/08482080A  
Patent No. 6010849  
GENERAL INFORMATION:  
APPLICANT: Edwards, Cynthia A.  
APPLICANT: Cantor, Charles R.  
APPLICANT: Andrews, Beth M.  
APPLICANT: Turin, Lisa M.

```

; APPLICANT: Fry, Kirk E.
; TITLE OF INVENTION: Sequence-Directed DNA Binding
; NUMBER OF SEQUENCES: 664
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Genelabs Technologies, Inc.
; STREET: 505 Penobscot Drive
; CITY: Redwood City
; STATE: CA
; COUNTRY: USA
; ZIP: 94063
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/482,080A
; FILING DATE: 07-JUN-1995
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/171,389
; FILING DATE: 20-DEC-1993
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/123,936
; FILING DATE: 17-SEP-1993
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/996,783
; FILING DATE: 23-DEC-1992
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/723,618
; FILING DATE: 27-JUN-1991
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/081,070
; FILING DATE: 22-JUN-1993
; ATTORNEY/AGENT INFORMATION:
; NAME: Brady, John F.
; REGISTRATION NUMBER: 39,118
; REFERENCE/DOCKET NUMBER: 4600-0175.20/G19P3D1
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (650) 324-0880
; TELEFAX: (650) 324-0960
; INFORMATION FOR SEQ ID NO: 546:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 50 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; HYPOTHETICAL: NO
; ORIGINAL SOURCE:
; INDIVIDUAL ISOLATE: Epstein Barr virus L2 (start site)
; INDIVIDUAL ISOLATE: 90021)
; US-08-482-080A-546

Query Match      88.9%; Score 16; DB 3; Length 50;
Best Local Similarity 100.0%; Pred. No. 1.2;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 GGCTGGTGCACCTGT 16
DB      41 GGCTGGTGCACCTGT 26

RESULT 6
US-09-354-947-546/c
; Sequence 546, Application US/09354947
; Patent No. 6384208
; GENERAL INFORMATION:
; APPLICANT: Edwards, Cynthia A.
; APPLICANT: Cantor, Charles R.
; APPLICANT: Andrews, Beth M.
; APPLICANT: Turin, Lisa M.
; APPLICANT: Fry, Kirk E.
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; TITLE OF INVENTION: Sequence-Directed DNA Binding
; NUMBER OF SEQUENCES: 664
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Genelabs Technologies, Inc.
; STREET: 505 Penobscot Drive
; CITY: Redwood City
; STATE: CA
; COUNTRY: USA
; ZIP: 94063
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/354,947
; FILING DATE:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/482,080
; FILING DATE: 07-JUN-1995
; APPLICATION NUMBER: US 08/171,389
; FILING DATE: 20-DEC-1993
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/123,936
; FILING DATE: 17-SEP-1993
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/996,783
; FILING DATE: 23-DEC-1992
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 07/723,618
; FILING DATE: 27-JUN-1991
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/081,070
; FILING DATE: 22-JUN-1993
; ATTORNEY/AGENT INFORMATION:
; NAME: Brady, John F.
; REGISTRATION NUMBER: 39,118
; REFERENCE/DOCKET NUMBER: 4600-0175.20/G19P3D1
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (650) 324-0880
; TELEFAX: (650) 324-0960
; INFORMATION FOR SEQ ID NO: 546:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 50 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; HYPOTHETICAL: NO
; ORIGINAL SOURCE:
; INDIVIDUAL ISOLATE: Epstein Barr virus L2 (start site)
; INDIVIDUAL ISOLATE: 90021)
; US-09-354-947-546

Query Match      88.9%; Score 16; DB 4; Length 50;
Best Local Similarity 100.0%; Pred. No. 1.2;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 GGCTGGTGCACCTGT 16
DB      41 GGCTGGTGCACCTGT 26

RESULT 7
PCT-US93-12388-546/c
; Sequence 546, Application PC/TUS9312388
; GENERAL INFORMATION:
; APPLICANT:
; TITLE OF INVENTION: Sequence-Directed DNA Binding
; NUMBER OF SEQUENCES: 641
; CORRESPONDENCE ADDRESS:
```



ADDRESSEE: Genelabs Technologies, Inc.  
STREET: 505 Penobscot Drive  
CITY: Redwood City  
STATE: CA  
COUNTRY: USA  
ZIP: 94063  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: Patent Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: PCT/US93/12388  
FILING DATE:  
CLASSIFICATION:  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/123,936  
FILING DATE: 17-SEP-1993  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/996,783  
FILING DATE: 23-DEC-1992  
ATTORNEY/AGENT INFORMATION:  
NAME: Fadian, Gary R.  
REGISTRATION NUMBER: 33,875  
REFERENCE/DOCKET NUMBER: 4600-0175.41/G19PCT2  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: (415) 324-0880  
TELEFAX: (415) 324-0960  
INFORMATION FOR SEQ ID NO: 546:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 50 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: double  
TOPOLOGY: linear  
MOLECULE TYPE: DNA (genomic)  
HYPOTHETICAL: NO  
ORIGINAL SOURCE:  
INDIVIDUAL ISOLATE: Epstein Barr virus L2 (start site  
INDIVIDUAL ISOLATE: 90021)  
PCT-US93-12388-546  
Query Match 88.9%; Score 16; DB 5; Length 50;  
Best Local Similarity 100.0%; Pred. No. 1.2;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 1 GGCTGGTGCACCTGT 16  
DB 41 GGCTGGTGCACCTGT 26  
RESULT 8  
US-08-917-320-18/c  
Sequence 18, Application US/08917320  
Patent No. 5824508  
GENERAL INFORMATION:  
APPLICANT: Spaete, Richard and Jackman, Winthrop, T.  
TITLE OF INVENTION: No. 5824508 Splicing Variants of gp350/220  
NUMBER OF SEQUENCES: 18  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Cooley Godward Castro Huddleson & Tatum  
STREET: 5 Palo Alto Square  
CITY: Palo Alto  
STATE: California  
COUNTRY: USA  
ZIP: 94306  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: Patent in Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/917,320  
FILING DATE: 25-AUG-1997

CLASSIFICATION: 435  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 08/229,291  
FILING DATE: April 18, 1994  
ATTORNEY/AGENT INFORMATION:  
NAME: Luann Caer  
REGISTRATION NUMBER: 31,822  
REFERENCE/DOCKET NUMBER: AVIR-003/00US  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 415-843-5163  
TELEFAX: 415-857-0663  
TELEX: 380816 CooleyPA  
INFORMATION FOR SEQ ID NO: 18:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 3833 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: double  
TOPOLOGY: unknown  
MOLECULE TYPE: CDNA  
FEATURE:  
NAME/KEY: CDS  
LOCATION: 1014..3734  
US-08-917-320-18  
Query Match 88.9%; Score 16; DB 1; Length 3833;  
Best Local Similarity 100.0%; Pred. No. 1;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 1 GGCTGGTGCACCTGT 16  
DB 3137 GGCTGGTGCACCTGT 3122  
RESULT 9  
PCT-US95-04611A-18/c  
Sequence 18, Application PC/TUS9504611A  
GENERAL INFORMATION:  
APPLICANT: Spaete, Richard and Jackman, Winthrop, T.  
TITLE OF INVENTION: Non Splicing Variants of gp350/220  
NUMBER OF SEQUENCES: 19  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Cooley Godward Castro Huddleson & Tatum  
STREET: 5 Palo Alto Square  
CITY: Palo Alto  
STATE: California  
COUNTRY: USA  
ZIP: 94306  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: Patent in Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: PCT/US95/04611A  
FILING DATE:  
CLASSIFICATION:  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 08/229,291  
FILING DATE: April 18, 1994  
ATTORNEY/AGENT INFORMATION:  
NAME: Luann Caer  
REGISTRATION NUMBER: 31,822  
REFERENCE/DOCKET NUMBER: AVIR-003/00US  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 415-843-5163  
TELEFAX: 415-857-0663  
TELEX: 380816 CooleyPA  
INFORMATION FOR SEQ ID NO: 18:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 3833 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: double

TOPOLOGY: unknown  
MOLECULE TYPE: CDNA  
FEATURE:  
NAME/KEY: CDS  
LOCATION: 1014..3734  
PCT-US95-04611A-18

Query Match 88.9%; Score 16; DB 5; Length 3833;  
Best Local Similarity 100.0%; Pred. No. 1;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGCTGGTGTACCTGT 16  
DB 3137 GGCTGGTGTACCTGT 3122

RESULT 10  
US-08-783-774-1/c

Sequence 1, Application US/08783774  
Patent No. 6054130  
GENERAL INFORMATION:  
APPLICANT: Spaete, Richard  
TITLE OF INVENTION: NON-SPLICING VARIANTS OF  
TITLE OF INVENTION: GP350/220  
NUMBER OF SEQUENCES: 19  
CORRESPONDENCE ADDRESSES:  
ADDRESSEE: Pennie & Edmonds  
STREET: 1155 Avenue of the Americas  
CITY: New York  
STATE: NY  
COUNTRY: USA  
ZIP: 10036/2711  
COMPUTER READABLE FORM:  
COMPUTER: IBM Compatible  
OPERATING SYSTEM: DOS  
SOFTWARE: FastSeq Version 2.0  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/783,774  
FILING DATE: 15-JAN-1997  
CLASSIFICATION: 435  
ATTORNEY/AGENT INFORMATION:  
NAME: Coruzzi, Laura A.  
REGISTRATION NUMBER: 30,742  
REFERENCE/DOCKET NUMBER: 7682-037  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 212-790-9090  
TELEFAX: 212-869-8864  
TELEX: 66141 PENNIE  
INFORMATION FOR SEQ ID NO: 1:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 5931 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
MOLECULE TYPE: DNA  
FEATURE:  
NAME/KEY: Coding Sequence  
LOCATION: 1014..3734  
OTHER INFORMATION:  
US-08-783-774-1

Query Match 88.9%; Score 16; DB 3; Length 5931;  
Best Local Similarity 100.0%; Pred. No. 0.99;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGCTGGTGTACCTGT 16  
DB 3137 GGCTGGTGTACCTGT 3122

RESULT 11

US-09-556-706B-1/c  
Sequence 1, Application US/09556706B  
Patent No. 6458364  
GENERAL INFORMATION:  
APPLICANT: Spaete, Richard  
TITLE OF INVENTION: NON-SPLICING VARIANTS OF GP350/220  
FILE REFERENCE: 7682-050-999  
CURRENT APPLICATION NUMBER: US/09/556,706B  
CURRENT FILING DATE: 2000-04-24  
PRIOR APPLICATION NUMBER: 08/783,774  
PRIOR FILING DATE: 1997-01-15  
PRIOR APPLICATION NUMBER: 08/229,291  
PRIOR FILING DATE: 1994-04-18  
NUMBER OF SEQ ID NOS: 19  
SOFTWARE: PatentIn version 3.0  
SEQ ID NO 1  
LENGTH: 5931  
TYPE: DNA  
ORGANISM: Virus  
FEATURE:  
OTHER INFORMATION: GP350/220  
US-09-556-706B-1

Query Match 88.9%; Score 16; DB 4; Length 5931;  
Best Local Similarity 100.0%; Pred. No. 0.99;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGCTGGTGTACCTGT 16  
DB 3137 GGCTGGTGTACCTGT 3122

RESULT 12  
US-09-171-710-5  
Sequence 5, Application US/09171710  
Patent No. 6323330  
GENERAL INFORMATION:  
APPLICANT: ISHIDUKA, Yasuyuki  
TITLE OF INVENTION: NOVEL PROTEINS C16 AND C16N OR GENES ENCODING THE SAME  
FILE REFERENCE: 0020-4474P  
CURRENT APPLICATION NUMBER: US/09/171,710  
CURRENT FILING DATE: 1998-10-23  
EARLIER APPLICATION NUMBER: PCT/JP97/01391  
EARLIER FILING DATE: 1997-10-30  
EARLIER APPLICATION NUMBER: 9-41562  
EARLIER FILING DATE: 1997-02-10  
EARLIER APPLICATION NUMBER: 8-127954  
EARLIER FILING DATE: 1996-04-23  
NUMBER OF SEQ ID NOS: 14  
SOFTWARE: PatentIn Ver. 2.0  
SEQ ID NO 5  
LENGTH: 1808  
TYPE: DNA  
ORGANISM: Homo sapiens  
US-09-171-710-5

Query Match 83.3%; Score 15; DB 4; Length 1808;  
Best Local Similarity 100.0%; Pred. No. 3.8;  
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGCTGGTGTACCTG 15  
DB 949 GGCTGGTGTACCTG 963

RESULT 13  
US-09-171-710-3  
Sequence 3, Application US/09171710  
Patent No. 6323330  
GENERAL INFORMATION:  
APPLICANT: ISHIDUKA, Yasuyuki

APPLICANT: MOCHIZUKI, Reiko  
TITLE OF INVENTION: NOVEL PROTEINS C16 AND C16N OR GENES ENCODING THE SAME  
FILE REFERENCE: 0020-4474P  
CURRENT APPLICATION NUMBER: US/09/171, 710  
CURRENT FILING DATE: 1998-10-23  
EARLIER APPLICATION NUMBER: PCT/JP97/01391  
EARLIER FILING DATE: 1997-10-30  
EARLIER APPLICATION NUMBER: 9-41562  
EARLIER FILING DATE: 1997-02-10  
EARLIER APPLICATION NUMBER: 8-127954  
EARLIER FILING DATE: 1996-04-23  
NUMBER OF SEQ ID NOS: 14  
SOFTWARE: PatentIn Ver. 2.0  
SEQ ID NO 3  
LENGTH: 3065  
TYPE: DNA  
ORGANISM: Mus sp.  
US-09-171-710-3

Query Match 83.3%; Score 15; DB 4; Length 3065;  
Best Local Similarity 100.0%; Pred. No. 3.7;  
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGCTGGTGCACCTG 15  
Db 1184 GGCTGGTGCACCTG 1198

RESULT 14  
US-07-745-206A-12  
Sequence 12, Application US/0745206A  
Patent No. 5429921  
GENERAL INFORMATION:  
APPLICANT: Harpold, Michael  
APPLICANT: Ellis, Steven  
APPLICANT: Williams, Mark  
APPLICANT: McCue, Ann  
APPLICANT: Feldman, Daniel  
TITLE OF INVENTION: Human Calcium Channel Compositions and  
TITLE OF INVENTION: Methods  
NUMBER OF SEQUENCES: 32  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Fitch, Even, Tabin & Flannery  
STREET: 135 S. Lasalle  
CITY: Chicago  
STATE: Illinois  
COUNTRY: U.S.A.  
ZIP: 60603  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: PatentIn Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/07/745,206A  
FILING DATE: 19910815  
CLASSIFICATION: 435  
ATTORNEY/AGENT INFORMATION:  
NAME: Feder, Scott B  
REFERENCE/DOCKET NUMBER: 51504  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 312-372-7842  
INFORMATION FOR SEQ ID NO: 12:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 5467 base pairs  
TYPE: NUCLEIC ACID  
STRANDEDNESS: unknown  
TOPOLOGY: unknown  
MOLECULE TYPE: DNA (genomic)  
FEATURE:  
NAME/KEY: CDS  
LOCATION: join(144..3164, 3168..3245, 3249..3386, 3390  
LOCATION: ..3392, 3396..3488, 3495..3539, 3543..3581, 3585

LOCATION: ..3587, 3591..3626, 3630..3689, 3693..3737, 3744  
LOCATION: ..3746, 3750..4823, 4827..4841, 4845..5006, 5010  
LOCATION: ..5096, 5100..5306, 5310..5366, 5370..5465)  
US-07-745-206A-12

Query Match 77.8%; Score 14; DB 1; Length 5467;  
Best Local Similarity 100.0%; Pred. No. 14;  
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 CTGGTGCACCTG 16  
Db 4965 CTGGTGCACCTG 4978

RESULT 15  
US-08-311-363-12  
Sequence 12, Application US/08311363  
Patent No. 5876958  
GENERAL INFORMATION:  
APPLICANT: Harpold, Michael  
APPLICANT: Ellis, Steven  
APPLICANT: Williams, Mark  
APPLICANT: Feldman, Daniel  
APPLICANT: McCue, Ann  
APPLICANT: Brenner, Robert  
TITLE OF INVENTION: Human Calcium Channel Compositions and  
TITLE OF INVENTION: Methods  
NUMBER OF SEQUENCES: 32  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Brown, Martin, Haller & McClain  
STREET: 1660 Union Street  
CITY: San Diego  
STATE: California  
COUNTRY: USA  
ZIP: 92101-2926  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: PatentIn Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/311,363  
FILING DATE:  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/745,206  
FILING DATE: 15-AUG-1991  
ATTORNEY/AGENT INFORMATION:  
NAME: Seidman, Stephanie L.  
REGISTRATION NUMBER: 33,779  
REFERENCE/DOCKET NUMBER: 6362-51506  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: (619)238-0999  
TELEFAX: (619)238-0062  
INFORMATION FOR SEQ ID NO: 12:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 5467 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: unknown  
TOPOLOGY: unknown  
MOLECULE TYPE: DNA (genomic)  
FEATURE:  
NAME/KEY: CDS  
LOCATION: join(144..3164, 3168..3245, 3249..3386, 3390  
LOCATION: ..3392, 3396..3488, 3495..3539, 3543..3581, 3585  
LOCATION: ..3587, 3591..3626, 3630..3689, 3693..3737, 3744  
LOCATION: ..3746, 3750..4823, 4827..4841, 4845..5006, 5010  
LOCATION: ..5096, 5100..5306, 5310..5366, 5370..5465)  
US-08-311-363-12

Query Match 77.8%; Score 14; DB 2; Length 5467;  
Best Local Similarity 100.0%; Pred. No. 14;  
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 3 CTGGTGTACCTGT 16  
Db 4965 CTGGTGTACCTGT 4978

RESULT 16  
US-09-657-453A-24/c  
Sequence 24, Application US/09657453A  
Patent No. 6458591  
GENERAL INFORMATION:  
APPLICANT: Brett P. Monia  
TITLE OF INVENTION: ANTISENSE MODULATION OF PHOSPHORYLASE KINASE ALPHA 2 EXPRESS  
FILE REFERENCE: RTS-0136  
CURRENT APPLICATION NUMBER: US/09/657,453A  
CURRENT FILING DATE: 2000-09-07  
NUMBER OF SEQ ID NOS: 105  
SEQ ID NO 24  
LENGTH: 261  
TYPE: DNA  
ORGANISM: Homo sapiens  
FEATURE:  
NAME/KEY: CDS  
LOCATION: (148)...(177)  
US-09-657-453A-24

Query Match 72.2%; Score 13; DB 4; Length 261;  
Best Local Similarity 100.0%; Pred. No. 56;  
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 5 GGTGTACCTGTT 17  
Db 185 GGTGTACCTGTT 173

RESULT 17  
US-09-141-027-9/c  
Sequence 9, Application US/09141027A  
Patent No. 6372454  
GENERAL INFORMATION:  
APPLICANT: Duan, et al.  
TITLE OF INVENTION: Follistatin-3  
FILE REFERENCE: PF388  
CURRENT APPLICATION NUMBER: US/09/141,027A  
CURRENT FILING DATE: 1998-08-27  
EARLIER APPLICATION NUMBER: 60/656,248  
EARLIER FILING DATE: 1997-08-29  
NUMBER OF SEQ ID NOS: 19  
SOFTWARE: Patentin Ver. 2.0  
SEQ ID NO 9  
LENGTH: 308  
TYPE: DNA  
ORGANISM: Homo sapiens  
FEATURE:  
NAME/KEY: misc\_feature  
LOCATION: (3)  
OTHER INFORMATION: n equals a, t, g, or c  
FEATURE:  
NAME/KEY: misc\_feature  
LOCATION: (19)  
OTHER INFORMATION: n equals a, t, g, or c  
FEATURE:  
NAME/KEY: misc\_feature  
LOCATION: (24)  
OTHER INFORMATION: n equals a, t, g, or c  
FEATURE:  
NAME/KEY: misc\_feature  
LOCATION: (29)  
OTHER INFORMATION: n equals a, t, g, or c  
FEATURE:  
NAME/KEY: misc\_feature  
LOCATION: (34)  
OTHER INFORMATION: n equals a, t, g, or c

FEATURE:  
NAME/KEY: misc\_feature  
LOCATION: (38)  
OTHER INFORMATION: n equals a, t, g, or c  
FEATURE:  
NAME/KEY: misc\_feature  
LOCATION: (40)  
OTHER INFORMATION: n equals a, t, g, or c  
FEATURE:  
NAME/KEY: misc\_feature  
LOCATION: (50)  
OTHER INFORMATION: n equals a, t, g, or c  
FEATURE:  
NAME/KEY: misc\_feature  
LOCATION: (83)  
OTHER INFORMATION: n equals a, t, g, or c  
FEATURE:  
NAME/KEY: misc\_feature  
LOCATION: (107)  
OTHER INFORMATION: n equals a, t, g, or c  
FEATURE:  
NAME/KEY: misc\_feature  
LOCATION: (205)  
OTHER INFORMATION: n equals a, t, g, or c  
FEATURE:  
NAME/KEY: misc\_feature  
LOCATION: (220)  
OTHER INFORMATION: n equals a, t, g, or c  
FEATURE:  
NAME/KEY: misc\_feature  
LOCATION: (237)  
OTHER INFORMATION: n equals a, t, g, or c  
FEATURE:  
NAME/KEY: misc\_feature  
LOCATION: (272)  
OTHER INFORMATION: n equals a, t, g, or c  
FEATURE:  
NAME/KEY: misc\_feature  
LOCATION: (297)  
OTHER INFORMATION: n equals a, t, g, or c  
FEATURE:  
NAME/KEY: misc\_feature  
LOCATION: (308)  
OTHER INFORMATION: n equals a, t, g, or c  
US-09-141-027-9

Query Match 72.2%; Score 13; DB 4; Length 308;  
Best Local Similarity 100.0%; Pred. No. 55;  
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 2 GGTGTGTACCT 14  
Db 18 GGTGTGTACCT 6

RESULT 18  
US-09-617-804-9/c  
Sequence 9, Application US/09617804  
Patent No. 6537966  
GENERAL INFORMATION:  
APPLICANT: Duan, et al.  
TITLE OF INVENTION: Follistatin-3  
FILE REFERENCE: PF388P1  
CURRENT APPLICATION NUMBER: US/09/617,804  
CURRENT FILING DATE: 2000-07-14  
PRIOR APPLICATION NUMBER: 60/144,088  
PRIOR FILING DATE: 1999-07-16  
PRIOR APPLICATION NUMBER: 09/141,027  
PRIOR FILING DATE: 1998-08-27

PRIOR APPLICATION NUMBER: 60/056,248  
PRIOR FILING DATE: 1997-08-29  
PRIOR APPLICATION NUMBER: PCT/US98/17710  
PRIOR FILING DATE: 1998-08-27  
NUMBER OF SEQ ID NOS: 22  
SOFTWARE: PatentIn Ver. 2.0  
SEQ ID NO 9  
LENGTH: 308  
TYPE: DNA  
ORGANISM: Homo sapiens  
FEATURE:  
NAME/KEY: misc\_feature  
LOCATION: (3)  
OTHER INFORMATION: n equals a, t, g, or c  
NAME/KEY: misc\_feature  
LOCATION: (19)  
OTHER INFORMATION: n equals a, t, g, or c  
NAME/KEY: misc\_feature  
LOCATION: (24)  
OTHER INFORMATION: n equals a, t, g, or c  
NAME/KEY: misc\_feature  
LOCATION: (29)  
OTHER INFORMATION: n equals a, t, g, or c  
NAME/KEY: misc\_feature  
LOCATION: (34)  
OTHER INFORMATION: n equals a, t, g, or c  
NAME/KEY: misc\_feature  
LOCATION: (38)  
OTHER INFORMATION: n equals a, t, g, or c  
NAME/KEY: misc\_feature  
LOCATION: (40)  
OTHER INFORMATION: n equals a, t, g, or c  
NAME/KEY: misc\_feature  
LOCATION: (50)  
OTHER INFORMATION: n equals a, t, g, or c  
NAME/KEY: misc\_feature  
LOCATION: (83)  
OTHER INFORMATION: n equals a, t, g, or c  
NAME/KEY: misc\_feature  
LOCATION: (107)  
OTHER INFORMATION: n equals a, t, g, or c  
NAME/KEY: misc\_feature  
LOCATION: (205)  
OTHER INFORMATION: n equals a, t, g, or c  
NAME/KEY: misc\_feature  
LOCATION: (220)  
OTHER INFORMATION: n equals a, t, g, or c  
NAME/KEY: misc\_feature  
LOCATION: (237)  
OTHER INFORMATION: n equals a, t, g, or c  
NAME/KEY: misc\_feature  
LOCATION: (272)  
OTHER INFORMATION: n equals a, t, g, or c  
NAME/KEY: misc\_feature  
LOCATION: (242)  
OTHER INFORMATION: n equals a, t, g, or c  
NAME/KEY: misc\_feature  
LOCATION: (297)  
OTHER INFORMATION: n equals a, t, g, or c  
NAME/KEY: misc\_feature  
LOCATION: (308)  
OTHER INFORMATION: n equals a, t, g, or c  
US-09-617-804-9

Query Match 72.2%; Score 13; DB 4; Length 308;  
Best Local Similarity 100.0%; Pred. No. 55;  
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 GCTGGTGTACCT 14  
DB 18 GCTGGTGTACCT 6

RESULT 19  
US-09-702-705-1159/c  
Sequence 1159, Application US/09702705  
Patent No. 6504010  
GENERAL INFORMATION:  
APPLICANT: Wang, Tongrong  
APPLICANT: Bangur, Chaitanya S.  
APPLICANT: Lodes, Michael A.  
APPLICANT: Fanger, Gary  
APPLICANT: Vedwick, Tom  
APPLICANT: Carter, Darrick  
APPLICANT: Retter, Marc  
APPLICANT: Mannion, Jane  
APPLICANT: Fan, Liqun  
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THE THERAPY AND  
FILE REFERENCE: 210121.478C14  
CURRENT APPLICATION NUMBER: US/09/702,705  
CURRENT FILING DATE: 2000-10-30  
NUMBER OF SEQ ID NOS: 1833  
SOFTWARE: FastSeq for Windows Version 3.0  
SEQ ID NO 1159  
LENGTH: 361  
TYPE: DNA  
ORGANISM: Homo sapien  
FEATURE:  
NAME/KEY: misc\_feature  
LOCATION: (1)...(361)  
OTHER INFORMATION: n = A,T,C or G  
US-09-702-705-1159

Query Match 72.2%; Score 13; DB 4; Length 361;  
Best Local Similarity 100.0%; Pred. No. 55;  
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 CTGGTGTACCTG 15  
DB 148 CTGGTGTACCTG 136

RESULT 20  
US-09-736-457-1159/c  
Sequence 1159, Application US/09736457  
Patent No. 6509448  
GENERAL INFORMATION:  
APPLICANT: Wang, Tongrong  
APPLICANT: Bangur, Chaitanya S.  
APPLICANT: Lodes, Michael A.  
APPLICANT: Fanger, Gary  
APPLICANT: Vedwick, Tom  
APPLICANT: Carter, Darrick  
APPLICANT: Retter, Marc  
APPLICANT: Mannion, Jane  
APPLICANT: Fan, Liqun  
APPLICANT: Wang, Aijun  
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THE THERAPY AND  
FILE REFERENCE: 210121.478C15  
CURRENT APPLICATION NUMBER: US/09/736,457  
CURRENT FILING DATE: 2000-12-13  
NUMBER OF SEQ ID NOS: 1864  
SOFTWARE: FastSeq for Windows Version 3.0  
SEQ ID NO 1159  
LENGTH: 361  
TYPE: DNA  
ORGANISM: Homo sapien  
FEATURE:  
NAME/KEY: misc\_feature  
LOCATION: (1)...(361)  
OTHER INFORMATION: n = A,T,C or G  
US-09-736-457-1159

Query Match 72.2%; Score 13; DB 4; Length 361;

Best Local Similarity 100.0%; Pred. No. 55;  
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 3 CTGGTGCACCTG 15  
|||||  
Db 148 CTGGTGCACCTG 136

## RESULT 21

US-09-228-986-40  
; Sequence 40, Application US/09228986  
; Patent No. 6359198  
; GENERAL INFORMATION:  
; APPLICANT: Strabala, Timothy  
; APPLICANT: Nieuwenhuizen, Niels  
; TITLE OF INVENTION: Compositions Isolated from Plant Cells  
; TITLE OF INVENTION: and their use in the Modification of Plant Cell Signalling  
; FILE REFERENCE: 11000/1020  
; CURRENT APPLICATION NUMBER: US/09/228,986  
; CURRENT FILING DATE: 1999-01-12  
; NUMBER OF SEQ ID NOS: 130  
; SOFTWARE: FastSeq for Windows Version 3.0  
; SEQ ID NO 40  
; LENGTH: 504  
; TYPE: DNA  
; ORGANISM: Pinus radiata  
US-09-228-986-40

Query Match 72.2%; Score 13; DB 4; Length 504;  
Best Local Similarity 100.0%; Pred. No. 54;  
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 3 CTGGTGCACCTG 15  
|||||  
Db 279 CTGGTGCACCTG 291

## RESULT 22

US-09-173-300-10  
; Sequence 10, Application US/09173300  
; Patent No. 6451581  
; GENERAL INFORMATION:  
; APPLICANT: Falco, Saverio Carl  
; APPLICANT: Hiltz, William D.  
; APPLICANT: Kinney, Anthony J.  
; APPLICANT: Cahoon, Rebecca E.  
; APPLICANT: Rafalski, J. Antoni  
; TITLE OF INVENTION: PLANT BRANCHED CHAIN AMINO ACID BIOSYNTHETIC ENZYMES  
; FILE REFERENCE: BB-1126  
; CURRENT APPLICATION NUMBER: US/09/173,300  
; CURRENT FILING DATE: 1998-10-15  
; EARLIER APPLICATION NUMBER: 60/063,423  
; EARLIER FILING DATE: 1997 October 28  
; NUMBER OF SEQ ID NOS: 54  
; SOFTWARE: Microsoft Word Version 7.0A  
; SEQ ID NO 10  
; LENGTH: 794  
; TYPE: DNA  
; ORGANISM: Zea mays  
US-09-173-300-10

Query Match 72.2%; Score 13; DB 4; Length 794;  
Best Local Similarity 100.0%; Pred. No. 54;  
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 4 TGGTGCACCTGT 16  
|||||  
Db 437 TGGTGCACCTGT 449

RESULT 23  
US-09-221-456-1/c  
; Sequence 1, Application US/09221456

; Patent No. 6162899  
; GENERAL INFORMATION:  
; APPLICANT: SATHE, GANESH  
; APPLICANT: HALSEY, WENDY  
; APPLICANT: MUIR, ALISON  
; APPLICANT: CHAMBERS, JON  
; APPLICANT: SZEKERES, PHILIP  
; TITLE OF INVENTION: METHODS OF SCREENING FOR AGONISTS  
; TITLE OF INVENTION: AND ANTAGONISTS OF THE HNEA81 RECEPTOR  
; NUMBER OF SEQUENCES: 2  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: Ratner & Prestia  
; STREET: P.O. Box 980  
; CITY: Valley Forge  
; STATE: PA  
; COUNTRY: USA  
; ZIP: 19482

COMPUTER READABLE FORM:

MEDIUM TYPE: Diskette

COMPUTER: IBM Compatible

OPERATING SYSTEM: DOS

SOFTWARE: FastSeq for Windows Version 2.0

CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/09/221,456

FILING DATE: 28-DEC-1998

CLASSIFICATION:

PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 08/956,975

FILING DATE: 23-OCT-1997

ATTORNEY/AGENT INFORMATION:  
NAME: Prestia, Paul F

REGISTRATION NUMBER: 23,031

REFERENCE/DOCKET NUMBER: GH-70318-1

TELECOMMUNICATION INFORMATION:  
TELEPHONE: 610-407-0700

TELEFAX: 610-407-0700

TELEX: 846169

INFORMATION FOR SEQ ID NO: 1:

SEQUENCE CHARACTERISTICS:

LENGTH: 1124 base pairs

TYPE: nucleic acid

STRANDEDNESS: single

TOPOLOGY: linear

MOLECULE TYPE: CDNA

US-09-221-456-1

Query Match 72.2%; Score 13; DB 3; Length 1124;  
Best Local Similarity 100.0%; Pred. No. 53;  
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 6 GTGTCACTGTGA 18  
|||||  
Db 89 GTGTCACTGTGA 77

## RESULT 24

US-09-558-740-1/c  
; Sequence 1, Application US/09558740  
; Patent No. 6358695  
; GENERAL INFORMATION:  
; APPLICANT: SATHE, GANESH  
; APPLICANT: HALSEY, WENDY  
; APPLICANT: MUIR, ALISON  
; APPLICANT: CHAMBERS, JON  
; APPLICANT: SZEKERES, PHILIP  
; TITLE OF INVENTION: METHODS OF SCREENING FOR AGONISTS AND  
; TITLE OF INVENTION: ANTAGONISTS OF THE HNEA81 RECEPTOR  
; FILE REFERENCE: GH-70318-2  
; CURRENT APPLICATION NUMBER: US/09/558,740  
; CURRENT FILING DATE: 2000-04-26  
; PRIOR APPLICATION NUMBER: 08/956,975  
; PRIOR FILING DATE: 1997-10-23  
; PRIOR APPLICATION NUMBER: 09/221,456

PRIOR FILING DATE: 1998-12-28  
NUMBER OF SEQ ID NOS: 2  
SOFTWARE: FastSeq for Windows Version 3.0  
SEQ ID NO 1  
LENGTH: 1124  
TYPE: DNA  
ORGANISM: HOMO SAPIENS  
US-09-558-740-1

Query Match 72.2%; Score 13; DB 4; Length 1124;  
Best Local Similarity 100.0%; Pred. No. 53;  
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 6 GTGTCACCTGTTA 18  
|||||  
DB 89 GTGTCACCTGTTA 77

RESULT 25  
US-09-328-352-3822/c  
Sequence 3822, Application US/09328352  
Patent No. 6562958  
GENERAL INFORMATION:  
APPLICANT: Gary L. Breton et al.  
TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO ACINETOBACTER  
FILE REFERENCE: GTC99-03PA  
CURRENT APPLICATION NUMBER: US/09/328,352  
FILING DATE: 1999-06-04  
NUMBER OF SEQ ID NOS: 8252  
SEQ ID NO 3822  
LENGTH: 1302  
TYPE: DNA  
ORGANISM: Acinetobacter baumannii  
US-09-328-352-3822

Query Match 72.2%; Score 13; DB 4; Length 1302;  
Best Local Similarity 100.0%; Pred. No. 53;  
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 TGGTGTACCTGT 16  
|||||  
DB 322 TGGTGTACCTGT 310

RESULT 26  
US-09-328-352-2907/c  
Sequence 2907, Application US/09328352  
Patent No. 6562958  
GENERAL INFORMATION:  
APPLICANT: Gary L. Breton et al.  
TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO ACINETOBACTER  
FILE REFERENCE: GTC99-03PA  
CURRENT APPLICATION NUMBER: US/09/328,352  
FILING DATE: 1999-06-04  
NUMBER OF SEQ ID NOS: 8252  
SEQ ID NO 2907  
LENGTH: 1716  
TYPE: DNA  
ORGANISM: Acinetobacter baumannii  
US-09-328-352-2907

Query Match 72.2%; Score 13; DB 4; Length 1716;  
Best Local Similarity 100.0%; Pred. No. 52;  
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 6 GTGTCACCTGTTA 18  
|||||  
DB 1328 GTGTCACCTGTTA 1316

RESULT 27

US-09-099-676-2  
Sequence 2, Application US/09099676  
Patent No. 6100075  
GENERAL INFORMATION:  
APPLICANT: Hillman, Jennifer L.  
APPLICANT: Corley, Neil C.  
APPLICANT: Baughn, Mariah R.  
TITLE OF INVENTION: DELTA 1-PYRROLINE-5-CARBOXYLATE REDUCTASE  
NUMBER OF SEQUENCES: 3  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Incyte Pharmaceuticals, Inc.  
STREET: 3174 Porter Drive  
CITY: Palo Alto  
STATE: CA  
COUNTRY: USA  
ZIP: 94304

COMPUTER READABLE FORM:  
MEDIUM TYPE: Diskette  
COMPUTER: IBM Compatible  
OPERATING SYSTEM: Windows  
SOFTWARE: FastSeq for Windows Version 2.0b  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/09/099,676  
FILING DATE: HERewith  
CLASSIFICATION:  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER:  
FILING DATE:  
ATTORNEY/AGENT INFORMATION:  
NAME: Cetrone, Michael C  
REGISTRATION NUMBER: 39,132  
REFERENCE/DOCKET NUMBER: PF-0532 US  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 650-855-0555  
TELEFAX: 650-855-0572  
TELEX:

INFORMATION FOR SEQ ID NO: 2:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 1742 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
IMMEDIATE SOURCE:  
LIBRARY: PROSONO1  
CLONE: 2278458  
US-09-099-676-2

Query Match 72.2%; Score 13; DB 3; Length 1742;  
Best Local Similarity 100.0%; Pred. No. 52;  
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGCTGTGTACCC 13  
|||||  
DB 479 GGCTGTGTACCC 491

RESULT 28  
US-09-565-910-2  
Sequence 2, Application US/09565910  
Patent No. 6268192  
GENERAL INFORMATION:  
APPLICANT: Hillman, Jennifer L.  
APPLICANT: Corley, Neil C.  
APPLICANT: Baughn, Mariah R.  
TITLE OF INVENTION: DELTA 1-PYRROLINE-5-CARBOXYLATE REDUCTASE  
NUMBER OF SEQUENCES: 3  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Incyte Pharmaceuticals, Inc.  
STREET: 3174 Porter Drive  
CITY: Palo Alto  
STATE: CA

COUNTRY: USA  
ZIP: 94304  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Diskette  
COMPUTER: IBM Compatible  
OPERATING SYSTEM: Windows  
SOFTWARE: FastSeq for Windows Version 2.0b  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/09/565,910  
FILING DATE:  
CLASSIFICATION:  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 09/099,676  
FILING DATE:  
ATTORNEY/AGENT INFORMATION:  
NAME: Cerrone, Michael C  
REGISTRATION NUMBER: 39,132  
REFERENCE/DOCKET NUMBER: PF-0532 US  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 650-855-0555  
TELEFAX: 650-855-0572  
TELEX:  
INFORMATION FOR SEQ ID NO: 2:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 1742 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
IMMEDIATE SOURCE:  
LIBRARY: PROSNON01  
CLONE: 2278458  
US-09-565-910-2

Query Match 72.2%; Score 13; DB 3; Length 1742;  
Best Local Similarity 100.0%; Pred. No. 52;  
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGCTGGTGCACC 13  
Db 479 GGCTGGTGCACC 491

RESULT 29  
US-08-454-455-3/C  
Sequence 3, Application US/08454455  
Patent No. 5635601  
GENERAL INFORMATION:  
APPLICANT: Moyle, Matthew  
APPLICANT: McLean, John W.  
TITLE OF INVENTION: NOVEL BETA INTEGRIN SUBUNIT  
NUMBER OF SEQUENCES: 9  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Genentech, Inc.  
STREET: 460 Point San Bruno Blvd  
CITY: South San Francisco  
STATE: California  
COUNTRY: USA  
ZIP: 94080  
COMPUTER READABLE FORM:  
MEDIUM TYPE: 3.5 inch, 720 Kb floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: WinPatIn (Genentech)  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/454,455  
FILING DATE: 30-May-1995  
CLASSIFICATION: 530  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 08/193989  
FILING DATE: 09-FEB-1994  
PRIOR APPLICATION DATA: 08/004142  
FILING DATE: 13-JAN-1993

PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 07/670607  
FILING DATE: 14-MAR-1991  
ATTORNEY/AGENT INFORMATION:  
NAME: Lee, Wendy M.  
REGISTRATION NUMBER: 00,000  
REFERENCE/DOCKET NUMBER: P0699C2D2  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 415/952-9881  
TELEFAX: 415/952-9881  
TELEX: 910/371-7168  
INFORMATION FOR SEQ ID NO: 3:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 2972 base pairs  
TYPE: Nucleic Acid  
STRANDEDNESS: Single  
TOPOLOGY: Linear  
US-08-454-455-3

Query Match 72.2%; Score 13; DB 1; Length 2972;  
Best Local Similarity 100.0%; Pred. No. 51;  
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 CTGGTGCACCTG 15  
Db 1012 CTGGTGCACCTG 1000

RESULT 30  
US-08-454-455-5/C  
Sequence 5, Application US/08454455  
Patent No. 5635601  
GENERAL INFORMATION:  
APPLICANT: Moyle, Matthew  
APPLICANT: McLean, John W.  
TITLE OF INVENTION: NOVEL BETA INTEGRIN SUBUNIT  
NUMBER OF SEQUENCES: 9  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Genentech, Inc.  
STREET: 460 Point San Bruno Blvd  
CITY: South San Francisco  
STATE: California  
COUNTRY: USA  
ZIP: 94080  
COMPUTER READABLE FORM:  
MEDIUM TYPE: 3.5 inch, 720 Kb floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: WinPatIn (Genentech)  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/454,455  
FILING DATE: 30-May-1995  
CLASSIFICATION: 530  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 08/193989  
FILING DATE: 09-FEB-1994  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 08/004142  
FILING DATE: 13-JAN-1993  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 07/670607  
FILING DATE: 14-MAR-1991  
ATTORNEY/AGENT INFORMATION:  
NAME: Lee, Wendy M.  
REGISTRATION NUMBER: 00,000  
REFERENCE/DOCKET NUMBER: P0699C2D2  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 415/952-9881  
TELEFAX: 415/952-9881  
TELEX: 910/371-7168  
INFORMATION FOR SEQ ID NO: 5:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 3789 base pairs



TYPE: Nucleic Acid  
STRANDEDNESS: Single  
TOPOLOGY: Linear  
US-08-454-455-5

Query Match 72.2%; Score 13; DB 1; Length 3789;  
Best Local Similarity 100.0%; Pred. No. 51;  
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 3 CTGGTGTACCTG 15  
|||||  
DB 1041 CTGGTGTACCTG 1029

RESULT 31  
US-08-474-067-1  
Sequence 1, Application US/08474067  
Patent No. 5811518  
GENERAL INFORMATION:  
APPLICANT: Ranscht, Barbara  
TITLE OF INVENTION: T-Cadherin Adhesion Molecule  
NUMBER OF SEQUENCES: 9  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Campbell and Flores  
STREET: 4370 La Jolla Village Drive, Suite 700  
CITY: San Diego  
STATE: California  
COUNTRY: United States  
ZIP: 92122  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: Patentin Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/474,067  
FILING DATE: 07-JUN-1995  
CLASSIFICATION: 514  
PRIORITY APPLICATION DATA:  
APPLICATION NUMBER: US 08/213,361  
FILING DATE: 14-MAY-1994  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/607,293  
FILING DATE: 30-OCT-1990  
ATTORNEY/AGENT INFORMATION:  
NAME: Campbell, Cathryn A.  
REGISTRATION NUMBER: 31,815  
REFERENCE/DOCKET NUMBER: P-LJ 1682  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: (619) 535-9001  
TELEFAX: (619) 535-8949  
INFORMATION FOR SEQ ID NO: 1:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 3959 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
FEATURE:  
NAME/KEY: CDS  
LOCATION: 45..2181  
US-08-474-067-1

Query Match 72.2%; Score 13; DB 1; Length 3959;  
Best Local Similarity 100.0%; Pred. No. 51;  
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 4 TGGTGTACCTGT 16  
|||||  
DB 976 TGGTGTACCTGT 988

RESULT 32  
US-08-474-068A-1

Sequence 1, Application US/08474068A

Patent No. 5837525

GENERAL INFORMATION:

APPLICANT: Ranscht, Barbara

TITLE OF INVENTION: T-Cadherin Adhesion Molecule

NUMBER OF SEQUENCES: 9

CORRESPONDENCE ADDRESS:

ADDRESSEE: Campbell & Flores LLP

STREET: 4370 La Jolla Village Drive, Suite 700

CITY: San Diego

STATE: California

COUNTRY: United States

ZIP: 92122

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk

COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: Patentin Release #1.0, Version #1.25

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/474,068A

FILING DATE: 07-JUN-1995

CLASSIFICATION: 514

PRIORITY APPLICATION DATA:

APPLICATION NUMBER: US 08/213,361

FILING DATE: 14-MAY-1994

PRIOR APPLICATION DATA:

APPLICATION NUMBER: US 07/607,293

FILING DATE: 30-OCT-1990

ATTORNEY/AGENT INFORMATION:

NAME: Campbell, Cathryn A.

REGISTRATION NUMBER: 31,815

REFERENCE/DOCKET NUMBER: P-LJ 1683

TELECOMMUNICATION INFORMATION:

TELEPHONE: (619) 535-9001

TELEFAX: (619) 535-8949

INFORMATION FOR SEQ ID NO: 1:

SEQUENCE CHARACTERISTICS:

LENGTH: 3959 base pairs

TYPE: nucleic acid

STRANDEDNESS: single

TOPOLOGY: linear

FEATURE:

NAME/KEY: CDS

LOCATION: 45..2181

US-08-474-068A-1

Query Match 72.2%; Score 13; DB 2; Length 3959;  
Best Local Similarity 100.0%; Pred. No. 51;  
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 4 TGGTGTACCTGT 16  
|||||  
DB 976 TGGTGTACCTGT 988

RESULT 33  
US-08-472-481-1

Sequence 1, Application US/08472481

Patent No. 5863804

GENERAL INFORMATION:

APPLICANT: Ranscht, Barbara

TITLE OF INVENTION: T-Cadherin Adhesion Molecule

NUMBER OF SEQUENCES: 8

CORRESPONDENCE ADDRESS:

ADDRESSEE: Campbell and Flores

STREET: 4370 La Jolla Village Drive, Suite 700

CITY: San Diego

STATE: California

COUNTRY: United States

ZIP: 92122

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk

COMPUTER: IBM PC compatible.

OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: PatentIn Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/472,481  
FILING DATE: 07-JUN-1995  
CLASSIFICATION: 424  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/213,361  
FILING DATE: 14-MAY-1994  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/607,293  
FILING DATE: 30-OCT-1990  
ATTORNEY/AGENT INFORMATION:  
NAME: Campbell, Cathryn A.  
REGISTRATION NUMBER: 31,815  
REFERENCE/DOCKET NUMBER: P-LJ 1686  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: (619) 535-9001  
TELEFAX: (619) 535-8949  
INFORMATION FOR SEQ ID NO: 1:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 3959 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
FEATURE:  
NAME/KEY: CDS  
LOCATION: 45..2181  
US-08-472-481-1

Query Match 72.2%; Score 13; DB 2; Length 3959;  
Best Local Similarity 100.0%; Pred. No. 51;  
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 TGCTGTCACTGT 16  
|||||  
Db 976 TGCTGTCACTGT 988

RESULT 34  
PCT-US96-01314-39/c  
Sequence 39, Application PC/TUS9601314  
GENERAL INFORMATION:  
APPLICANT: M. Amin Arnaut  
TITLE OF INVENTION: METHODS FOR IDENTIFYING INTEGRIN  
NUMBER OF SEQUENCES: 78  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Fish & Richardson P.C.  
STREET: 225 Franklin Street  
CITY: Boston  
STATE: Massachusetts  
COUNTRY: U.S.A.  
ZIP: 02110-2804  
COMPUTER READABLE FORM:  
MEDIUM TYPE: 3.5" Diskette, 1.44 Mb  
COMPUTER: IBM PS/2 Model 50Z or 55SX  
OPERATING SYSTEM: MS-DOS (Version 5.0)  
SOFTWARE: WordPerfect (Version 5.1)  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: PCT/US96/01314  
FILING DATE: 30-JAN-96  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 08/380,167  
FILING DATE: 30-JAN-95  
ATTORNEY/AGENT INFORMATION:  
NAME: John W. Freeman  
REGISTRATION NUMBER: 29,066  
REFERENCE/DOCKET NUMBER: 00786/267001  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: (617) 542-5070  
TELEFAX: (617) 542-8906  
TELEX: 200154

INFORMATION FOR SEQ ID NO: 39:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 5137 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
MOLECULE TYPE: DNA (genomic)  
PCT-US96-01314-39

Query Match 72.2%; Score 13; DB 5; Length 5137;  
Best Local Similarity 100.0%; Pred. No. 50;  
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGCTGGTGCACC 13  
|||||  
Db 4208 GGCTGGTGCACC 4196

RESULT 35  
US-08-476-062A-39/c  
Sequence 39, Application US/08476062A  
Patent No. 5877275  
GENERAL INFORMATION:  
APPLICANT: Arnaut, M. Amin  
TITLE OF INVENTION: CONTROLLING CELLULAR IMMUNE/INFLAMMATORY  
NUMBER OF SEQUENCES: 53  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Fish & Richardson P.C.  
STREET: 225 Franklin Street  
CITY: Boston  
STATE: MA  
COUNTRY: US  
ZIP: 02110-2804  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Diskette  
COMPUTER: IBM Compatible  
OPERATING SYSTEM: Windows95  
SOFTWARE: FastSeq for Windows Version 2.0  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/476,062A  
FILING DATE: 07-JUN-1995  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 08/216,081  
FILING DATE: 21-MAR-1994  
APPLICATION NUMBER: 07/637,830  
FILING DATE: 04-JAN-1991  
APPLICATION NUMBER: 07/539,842  
FILING DATE: 18-JUN-1990  
APPLICATION NUMBER: 07/212,573  
FILING DATE: 28-JUN-1988  
ATTORNEY/AGENT INFORMATION:  
NAME: Freeman, John W.  
REGISTRATION NUMBER: 29,066  
REFERENCE/DOCKET NUMBER: 00786/068003  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 617/542-5070  
TELEFAX: 617/542-8906  
TELEX: 200154  
INFORMATION FOR SEQ ID NO: 39:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 5138 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
MOLECULE TYPE: cDNA  
FEATURE:  
NAME/KEY: Coding Sequence  
LOCATION: 95..3604  
US-08-476-062A-39

Query Match 72.2%; Score 13; DB 2; Length 5138;  
Best Local Similarity 100.0%; Pred. No. 50;

Mon Aug 18 10:30:06 2003

us-10-074-620-1.011.rn1

Page 15

Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GGCTGGTGCACC 13

Db 4209 GGCTGGTGCACC 4197

Search completed: August 15, 2003, 11:00:16  
Job time : 43.025 secs

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GenCore version 5.1.6  
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OM nucleic - nucleic search, using sw model

Run on: August 14, 2003, 20:57:44 ; Search time 113.4 Seconds  
(without alignments)  
428.482 Million cell updates/sec

Title: US-10-074-620-1  
Perfect score: 18  
Sequence: 1 ggcgtgctgcacctgttca 18

Scoring table: OLIGO\_NUC  
Gapop 60.0 , Gapext 60.0

Searched: 2552756 seqs, 1349719017 residues

Word size: 0

Total number of hits satisfying chosen parameters: 5105512

Minimum DB seq length: 0  
Maximum DB seq length: 2000000000

Post-processing: Listing first 120 summaries

Database: N\_Geneseq\_19Jun03.\*

- 1: /SIDSI/gcgdata/geneq/geneq-emb1/NA1980.DAT.\*
- 2: /SIDSI/gcgdata/geneq/geneq-emb1/NA1981.DAT.\*
- 3: /SIDSI/gcgdata/geneq/geneq-emb1/NA1982.DAT.\*
- 4: /SIDSI/gcgdata/geneq/geneq-emb1/NA1983.DAT.\*
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- 9: /SIDSI/gcgdata/geneq/geneq-emb1/NA1988.DAT.\*
- 10: /SIDSI/gcgdata/geneq/geneq-emb1/NA1989.DAT.\*
- 11: /SIDSI/gcgdata/geneq/geneq-emb1/NA1990.DAT.\*
- 12: /SIDSI/gcgdata/geneq/geneq-emb1/NA1991.DAT.\*
- 13: /SIDSI/gcgdata/geneq/geneq-emb1/NA1992.DAT.\*
- 14: /SIDSI/gcgdata/geneq/geneq-emb1/NA1993.DAT.\*
- 15: /SIDSI/gcgdata/geneq/geneq-emb1/NA1994.DAT.\*
- 16: /SIDSI/gcgdata/geneq/geneq-emb1/NA1995.DAT.\*
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- 19: /SIDSI/gcgdata/geneq/geneq-emb1/NA1998.DAT.\*
- 20: /SIDSI/gcgdata/geneq/geneq-emb1/NA1999.DAT.\*
- 21: /SIDSI/gcgdata/geneq/geneq-emb1/NA2000.DAT.\*
- 22: /SIDSI/gcgdata/geneq/geneq-emb1/NA2001.DAT.\*
- 23: /SIDSI/gcgdata/geneq/geneq-emb1/NA2002.DAT.\*
- 24: /SIDSI/gcgdata/geneq/geneq-emb1/NA2003.DAT.\*
- 25: /SIDSI/gcgdata/geneq/geneq-emb1/NA2004.DAT.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

# SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	100.0	18	24	ABA00268	EBNA-2 primer, Dos
2	88.9	20	16	AA091011	Primer binding to Epstein Barr virus
3	88.9	50	15	AA069796	Primer binding to EBV L2 (start site
4	88.9	50	18	AA064258	Test sequence from DNA binding molecu
5	88.9	50	24	ABX83037	Human secreted pro
6	88.9	435	21	AA001776	Human small cell ca
7	88.9	455	24	AA061522	
8	88.9	455	24	AA061522	

9	16	88.9	675	22	AA189104	Human polynucleoti
10	16	88.9	1142	19	AAV43611	Human secreted pro
11	16	88.9	1152	19	AAV62754	Human secreted pro
12	16	88.9	1152	24	ABO92057	Human polynucleoti
13	16	88.9	1217	21	AA077733	Human cancer assoc
14	16	88.9	1278	20	AAZ17770	Human gene expres
15	16	88.9	2382	25	ACC46555	Human dltmp secret
16	16	88.9	2921	6	AA050114	DNA sequence encod
17	16	88.9	5331	16	AA004821	EBV gp350/220 CDNA
18	16	88.9	13559	22	AA015144	Human nervous syst
19	15	83.3	65	24	ABN52794	Mouse spliced tran
20	15	83.3	609	23	ABV56544	Human prostate exp
21	15	83.3	616	21	AA011311	Aspergillus niger
22	15	83.3	1419	22	AAH45451	Murine epilepsy-ca
23	15	83.3	1699	24	AB211825	Human polynucleoti
24	15	83.3	1808	18	AAV02313	Clf6 gene for prom
25	15	83.3	2293	21	AAZ44730	Human Clf6N-1 cDNA
26	15	83.3	2293	21	AAZ40179	Human Clf6N-1 cDNA
27	15	83.3	2301	21	AAZ44731	Human Clf6N-2 cDNA
28	15	83.3	2301	21	AAZ40180	Human Clf6N-2 cDNA
29	15	83.3	2952	20	AAZ33584	Human breast tumo
30	15	83.3	3035	21	AA077831	Human cancer assoc
31	15	83.3	3035	22	AAH34609	Human colon cancer
32	15	83.3	3065	18	AAV02312	Clf6 gene for prom
33	15	83.3	3337	21	AAZ44729	Murine Clf6N-2 cDNA
34	15	83.3	3337	21	AAZ40178	Mouse Clf6N-2 codin
35	15	83.3	3674	21	AAZ44728	Murine Clf6N-1 cDNA
36	15	83.3	3674	21	AAZ40177	Mouse Clf6N-1 codin
37	15	83.3	4537	25	ABX63337	Human CDNA #337 di
38	15	83.3	6501	22	AA066882	Cocillibolus heter
39	15	83.3	6550	24	AB068409	Fungal peptide syn
40	15	83.3	6553	24	AB068449	DNA encoding C. he
41	15	83.3	6672	24	AB068121	Ovary cancer relat
42	15	83.3	7702	21	AA088739	Human protein tyro
43	15	83.3	7702	21	AA029198	Lar tyrosine phosph
44	15	83.3	7705	22	AA098405	Human EST-derived
45	15	83.3	7705	22	AA098405	Human EST-derived
46	15	83.3	7705	22	AA098405	Human EST-derived
47	15	83.3	7705	22	AA098405	Human EST-derived
48	15	83.3	7945	23	ABY27897	Human CDNA encodin
49	15	83.3	16831	23	ABY59607	Human prostate exp
50	15	83.3	42115	24	AB068452	Protonobacterium
51	14	77.8	249	25	AB072979	C. heterotrophus
52	14	77.8	327	22	AA180980	Rice leaf EST, SEQ
53	14	77.8	370	24	AB180865	Human polynucleoti
54	14	77.8	383	22	AAH99005	Human ovarian can
55	14	77.8	411	25	ABX50538	Murine EST-derived
56	14	77.8	454	22	AA072255	Bovine EST associa
57	14	77.8	455	22	AAH81534	DNA encoding novel
58	14	77.8	465	22	ABX52087	Human differential
59	14	77.8	465	22	ABX21899	Human foetal liver
60	14	77.8	465	22	AAK00369	Probe #365 for gen
61	14	77.8	465	22	AAK25812	Human brain expres
62	14	77.8	465	22	AAK25812	Human bone marrow
63	14	77.8	465	22	AAI31694	Probe #374 for gen
64	14	77.8	465	22	AAI31694	Probe #380 used to
65	14	77.8	465	22	AAI00377	Probe #368 used to
66	14	77.8	465	23	AB052400	Human liver single
67	14	77.8	465	23	AB050393	Human genome-deiv
68	14	77.8	1109	24	ABK35284	Human CDNA encodin
69	14	77.8	1220	22	AA046203	Human DNA encodin
70	14	77.8	1220	25	ACAS7961	Human PRO19626 CDN
71	14	77.8	1220	25	ABX98431	Human CDNA encodin
72	14	77.8	1220	25	ABX98933	Human CDNA encodin
73	14	77.8	1220	25	ABX98022	Novel human secret
74	14	77.8	1220	25	ABX98022	Human secreted/tra
75	14	77.8	1220	25	ABX78806	Human PRO polynucl
76	14	77.8	1220	25	ABX75819	Human CDNA encodin
77	14	77.8	1220	25	ABX77024	Human CDNA encodin
78	14	77.8	1220	25	ABX16864	Human CDNA encodin
79	14	77.8	1548	21	AA043772	Zea mays DNA fragm
80	14	77.8	2017	22	AB057811	Human CDNA encodin
81	14	77.8	2970	22	AAH99525	Human protein enco
					AA047350	Human transporter

C 82	14	77.8	3069	24	AAD36308	Human transporex
C 83	14	77.8	3088	24	ABZ35367	Human gene express
C 84	14	77.8	3913	22	AAH17826	Human cDNA sequenc
C 85	14	77.8	4447	23	AAST72881	DNA encoding novel
C 86	14	77.8	4614	23	AAST72559	DNA encoding novel
C 87	14	77.8	19628	22	AAK71839	Human immune/haema
C 88	14	77.8	32107	22	AAST32249	Human DNA repair a
C 89	14	77.8	32187	24	ABK567552	Novel Human DNA re
C 90	14	77.8	58837	24	ABK52612	Human Claspin geno
C 91	14	77.8	122888	24	ABK83569	Human cDNA diftere
C 92	13	72.2	60	24	ABN407553	Human spliced tian
C 93	13	72.2	145	23	AAAS58139	cDNA #815 encoding
C 94	13	72.2	261	24	ABK65127	Human phosphotylas
C 95	13	72.2	294	21	AAC25263	Human secreted pro
C 96	13	72.2	300	20	AAZ14789	Human gene express
C 97	13	72.2	308	20	AAZ28130	Human follistatin
C 98	13	72.2	308	22	AAD02679	HAOG52R cDNA clon
C 99	13	72.2	350	22	AAK61553	Human immune/haema
C 100	13	72.2	352	22	AAK76174	Human immune/haema
C 101	13	72.2	352	22	AAK76175	Human immune/haema
C 102	13	72.2	361	24	ABK39121	cDNA encoding Lung
C 103	13	72.2	361	25	ACA11450	Human Lung adenoca
C 104	13	72.2	361	25	ACA02636	Lung cancer therap
C 105	13	72.2	369	14	AAO50915	Human brain Exper
C 106	13	72.2	371	24	ABK29788	Colon adenocarcino
C 107	13	72.2	375	22	AAK35637	Human cardiovascul
C 108	13	72.2	375	22	AAK36753	Human cardiovascul
C 109	13	72.2	378	23	AAAS58045	cDNA #721 encoding
C 110	13	72.2	413	23	AAAS25389	Human ovarian PCR-
C 111	13	72.2	414	21	AAAG28247	Human colon cancer
C 112	13	72.2	445	23	AAST74105	DNA encoding novel
C 113	13	72.2	460	23	AAAS26480	DNA encoding novel
C 114	13	72.2	495	22	AAAS25208	Human ovarian PCR-
C 115	13	72.2	495	22	AAH83855	Human ovarian tumo
C 116	13	72.2	504	21	AAAT79302	Pinus radiata celli
C 117	13	72.2	506	22	AAAS24619	Human ovarian PCR-
C 118	13	72.2	506	22	AAH83182	Human ovarian tumo
C 119	13	72.2	527	22	AAH05185	Human reproductive
C 120	13	72.2	527	23	ABL968070	Human testicular a

## ALIGNMENTS

```

RESULT 1
ABAA00268
ID ABA00268 standard; DNA; 18 BP.
XX
XX
AC ABA00268;
XX
DT 29-NOV-2002 (first entry)
XX
DE EBNA 2 primer, Position 90030:90049.
XX
KM Primer; amplify; PCR; probe; detection; Epstein-Barr virus; EBV; ss
XX
OS Epstein-Barr virus.
XX
PN MO200264842-A2.
XX
PN 22-AUG-2002.
XX
PD 13-FEB-2002; 2002MO-US04339.
XX
PF 13-FEB-2002; 2002MO-US04339.
XX
PR 13-FEB-2001; 2001US-268439P.
XX
PA (CHIL-) CHILDRENS HOSPITAL RES FOUND.
XX
PI Witte DP, Groen PA;
XX
DR WPI: 2002-667015/71.
XX
XX New compositions comprising nucleic acid sequences which specifically

```

PT hybridizes to Epstein-Barr virus (EBV) nucleic acid, for detecting EBV  
 PT in clinical specimens to determine patients at high risk of to  
 PT developing EBV infections  
 PS  
 PS Claim 1; Page 44; 59pp; English.  
 XX  
 CC The sequences given in ABA00268-75 are primers and probes which were  
 CC used in the compositions of the invention for the detection of  
 CC Epstein-Barr virus (EBV). The compositions comprise at least one  
 CC purified and isolated oligonucleotide consisting of a nucleic acid  
 CC sequence which complements and specifically hybridizes to EBV nucleic  
 CC acid. The oligonucleotide sequences and compositions comprising them  
 CC are useful for detecting EBV in clinical specimens to determine  
 CC patients who are at high risk to develop serious and costly medical  
 CC complications, and allow for better clinical management of these  
 CC patients by earlier recognition of their infection status. The  
 CC oligonucleotide sequences may also be used to amplify EBV DNA  
 CC sequences. The use of the oligonucleotide sequences in the assay for  
 CC detecting EBV has a broad dynamic range of detection from less than  
 CC 10-100000000 copies of EBV DNA, is less labour intensive requiring only  
 CC one reaction tube for the EBV determination, highly sensitive, accurate  
 CC and has a rapid turn around time with assays that are completed,  
 CC including amplification, probe specific hybridization, and calculation  
 CC of copy number in less than 1 hour. The method may be adapted to  
 CC automated systems.  
 CC  
 SQ Sequence 18 BP; 2 A; 4 C; 6 G; 6 T; 0 other;  
 OY Query Match 100.0%; Score 18; DB 24; Length 18;  
 DB Best Local Similarity 100.0%; Pred. No. 0.41;  
 DB 1 GGCTGGTGTCACCTGTTA 18  
 1 GGCTGGTGTCACCTGTTA 18  
 RESULT 2  
 AA091011  
 ID AA091011 standard; DNA; 20 BP.  
 XX  
 XX AA091011;  
 AC  
 XX  
 XX 01-FEB-1996 (first entry)  
 DT  
 XX  
 XX Primer binding to 5' end of EBV nuc antigen gene.  
 DE  
 XX  
 XX Primer; PCR; amplification; DNA polymerase; exonuclease; Pfu; Taq;  
 KW Klenow fragment; T4; T7; Deep Vent; synthesis; mismatch; human; antibody;  
 KW heavy chain variable region; ss.  
 XX  
 XX Synthetic.  
 OS  
 XX  
 XX WO9516028-A1  
 BN  
 XX 15-JUN-1995.  
 PD  
 XX  
 XX 07-DEC-1994; 94WO-US14065.  
 PF  
 XX  
 XX 16-FEB-1994; 94US-0197791.  
 PR 08-DEC-1993; 93US-0164290.  
 PA (STRA-) STRATAGEME.  
 XX  
 XX Mullinax RL, Sorge JA;  
 PI  
 XX WPI; 1995-224316/29.  
 DR  
 XX  
 XX Compsn. useful for polynucleotide synthesis and cyclical  
 PT amplification - comprising a mixt. contg. an enzyme with 3'-5'  
 PT exonuclease activity and a DNA polymerase with less 3'-5'  
 PT exonuclease activity than the enzyme  
 XX

PS Examples; Page 35; 66pp; English.

CC Primers AA090984-091028 are examples of primers for testing a novel  
 CC composition for polynucleotide synthesis comprising a DNA polymerase  
 CC with high 3'-5' exonuclease activity in conjunction with a DNA polymerase  
 CC respectively. Other DNA polymerases containing high 3'-5' exonuclease  
 CC activity include E.coli DNA polymerase I, Klenow fragment, T4, T7, Vent  
 CC or Deep Vent DNA polymerases. The use of a DNA polymerase with high  
 CC 3'-5' exonuclease activity is designed to overcome the inability of DNA  
 CC polymerases with low 3'-5' exonuclease activities to initiate synthesis  
 CC from primers containing 3' terminal mismatches, e.g. due to errors  
 CC introduced during a PCR cycle.

CC This primer binds to a region at the 5' end of the Epstein-Barr  
 CC virus nuc antigen gene.

XX Sequence 20 BP; 2 A; 4 C; 7 G; 7 T; 0 other;

SO Query Match 88.9%; Score 16; DB 16; Length 20;  
 Best Local Similarity 100.0%; Pred. No. 5.8; Mismatches 0; Indels 0; Gaps 0;  
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 GGCTGGTGCACCTGT 16  
 DB 1 GGCTGGTGCACCTGT 16

RESULT 3  
 AA069796/c  
 ID AA069796 standard; DNA; 50 BP.

XX AA069796;  
 AC 25-MAR-2003 (updated)  
 DT 06-MAR-1995 (first entry)  
 XX 25-MAR-2003 (first entry)  
 DT 06-MAR-1995 (first entry)  
 XX Epstein Barr virus R1 L2 (start site 90021), target region.

DE DNA protein-binding assay; test sequence; screening sequence;  
 KM promoter; target; TATA box; Herpes Simplex Virus; HSV;  
 KM origin of replication; UL9; transcription factor; TFIID; ds.  
 XX Synthetic.

OS WO9414980-A1.  
 PN 07-JUL-1994.  
 PD 20-DEC-1993; 93WO-US12388.  
 XX 23-DEC-1992; 92US-0996783.  
 PR 17-SEP-1993; 93US-0123936.  
 XX (GENE-) GENELABS TECHNOLOGIES INC.

PA Andrews BM, Cantor CR, Edwards CA, Fry KE, Turin LM;  
 PI WPI; 1994-234711/28.  
 DR Sequence-directed DNA-binding molecules - useful in  
 XX pharmaceuticals and as molecular reagents  
 PT Claim 28; Page 485; 587pp; English.

XX A DNA protein-binding assay is provided, useful for screening  
 CC libraries of synthetic or biological cpds. for their ability  
 CC to bind DNA test sequences. The assay is versatile in that any  
 CC number of test sequences can be tested by placing the test sequence  
 CC adjacent to a defined protein-binding screening sequence. Binding  
 CC of moIs. to these test sequences changes the binding characteristics  
 CC of the protein mol. to its cognate binding sequence. When such a mol.  
 CC binds the test sequence, the equilibrium of the DNA:protein complexes  
 CC is disturbed, generating changes in the concentration of free DNA probe.

CC One application of this method is to eucaryotic general transcription  
 CC factors (e.g. TFIID), where the target region is typically selected  
 CC from DNA sequences adjacent to the binding site for the eucaryotic  
 CC transcription factor. Numerous exemplary test sequences are given:  
 CC the sequences in AA069251-731 and AA069850 correspond to promoter  
 CC targets (typically, TATA box-contg. sites) for human genes and the  
 CC sequences in AA069732-849 correspond to promoter targets for viral genes.  
 CC The test sequences may also be randomly generated. DNA:protein  
 CC interaction may be used for screening purposes, e.g. the Herpes Simplex  
 CC virus (HSV) origin of replication and UL9 (see AA069851-52, AA069865 and  
 CC AA069891).

XX (Updated on 25-MAR-2003 to correct PN field.)

SO Sequence 50 BP; 18 A; 14 C; 10 G; 8 T; 0 other;

XX Query Match 88.9%; Score 16; DB 15; Length 50;  
 Best Local Similarity 100.0%; Pred. No. 5.7; Mismatches 0; Indels 0; Gaps 0;  
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 GGCTGGTGCACCTGT 16  
 DB 41 GGCTGGTGCACCTGT 26

RESULT 4  
 AAT64258/c  
 ID AAT64258 standard; DNA; 50 BP.

XX AAT64258;  
 AC 25-MAR-2003 (updated)  
 DT 17-MAR-1997 (first entry)  
 XX EBV L2 (start site 90021) TFIID binding site.

DE Duplex DNA; target region; binding characteristic; DNA binding protein;  
 KM TFIID; transcription factor; binding site; inhibition; enhance;  
 KM cancer; inherited genetic disorder; ds.  
 XX Epstein-barr virus.

OS US5578444-A.  
 PN 26-NOV-1996.  
 PD 20-DEC-1993; 93US-0171389.  
 XX 20-DEC-1993; 93US-0171389.  
 PR 27-JUN-1991; 91US-0723618.  
 PR 23-DEC-1992; 92US-0996783.  
 PR 17-SEP-1993; 93US-0123936.  
 XX (GENE-) GENELABS TECHNOLOGIES INC.

PA Andrews BM, Cantor CR, Edwards CA, Fry KE, Turin LM;  
 PI WPI; 1997-020402/02.  
 DR Altering binding characteristics of DNA binding proteins to duplex  
 XX DNA - by attaching specific small cpd. to target region close to the  
 PT protein's binding site, useful in treatment of viral disease, cancer  
 PT etc

XX Claim 6; Column 377; 264pp; English.

XX The sequences given in AAT63713-4312 represent duplex DNA's which act  
 CC as target regions in the method of the invention. The method for  
 CC altering the binding characteristics of a DNA-binding protein to duplex  
 CC DNA comprises contacting the duplex DNA with a small molecule which  
 CC binds sequence-specifically to a target region, where, when the small  
 CC molecule is bound to the target region, it is adjacent to, but not  
 CC overlapping by more than 4 bp, a binding site for a DNA-binding protein.  
 CC The small molecule is added at a concentration effective to alter the

CC binding of the DNA binding protein, pref. TFIID, to its binding site on  
 CC the duplex DNA. The binding of the small molecule may inhibit or  
 CC enhance the binding of the DNA-binding protein to its binding site. The  
 CC compounds isolated using this method are potentially useful as  
 CC therapeutic agents for treatment of any disease which involves a  
 CC specific DNA sequence, e.g. cancer, or inherited genetic disorders etc.  
 CC The method is suitable for screening large biological or chemical  
 CC libraries and allows determination of sequence-specific and relative  
 CC affinities of known DNA-binding agents for different DNA sequences.  
 CC The design of these duplex DNA's allows a single DNA-protein interaction  
 CC to be used for screening sequence-specific, or preferential, DNA binding  
 CC proteins that recognise almost any possible sequence (see also AAT49539-  
 CC 74).

CC (Updated on 25-MAR-2003 to correct PF field.)

XX  
 CC  
 SQ Sequence 50 BP; 18 A; 14 C; 10 G; 8 T; 0 other;

Query Match 88.9%; Score 16; DB 18; Length 50;  
 Best Local Similarity 100.0%; Pred. No. 5.7;  
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GGCTGCTGCACCTGT 16  
 Db 41 GGCTGCTGCACCTGT 26

RESULT 5  
 AAX17546/C  
 ID AAX17546 standard; DNA; 50 BP.  
 XX  
 AC AAX17546;  
 XX  
 DT 06-MAY-1999 (first entry)  
 XX  
 DE Test sequence from Epstein Barr virus L2 (start site 90021).  
 XX  
 KM Test sequence; DNA-binding molecule; screening sequence; human;  
 KM nucleic acid amplification; target; viral; ds.  
 XX  
 OS Epstein-Barr virus.  
 OS  
 PN US5869241-A.  
 PN  
 PD 09-FEB-1999.  
 PD  
 PF 07-JUN-1995; 95US-0475228.  
 PF  
 XX 20-DEC-1993; 93US-0171389.  
 PR 27-JUN-1991; 91US-0723618.  
 PR 23-DEC-1992; 92US-0896783.  
 PR 17-SEP-1993; 93US-0123936.  
 PR 07-JUN-1995; 95US-0475228.  
 PR  
 XX  
 PA (GENE-) GENELABS TECHNOLOGIES INC.  
 PA  
 PI Andrews BM, Cantor CR, Edwards CA, Fry KE, Turin LM;  
 PI  
 XX WPI; 1999-152755/13.  
 DR  
 XX  
 PT Determination of DNA sequence preference of a DNA-binding molecule -  
 PT based on inhibition of binding of protein to oligonucleotide  
 PT sequence attached to test sequence  
 PT  
 XX  
 PS Claim 3; Columns 379-380; 270pp; English.  
 PS  
 XX  
 CC Sequences AAX17001 to AAX17600 represent specifically claimed target  
 CC test sequences that are used in the method of the invention of  
 CC determining the DNA sequence preference of a DNA-binding molecule. The  
 CC method comprises: (i) adding a test molecule and a DNA-binding protein to  
 CC a mixture of duplex DNA test oligonucleotides, each of the test  
 CC oligonucleotides having a test sequence adjacent to a screening sequence,  
 CC where the screening sequence binds to the DNA-binding protein with a  
 CC binding affinity that is independent of the DNA sequence of the test

CC sequence, and where the mixture of duplex DNA test oligonucleotides  
 CC includes several test sequences; (ii) incubating the test molecule, the  
 CC mixture of duplex DNA test oligonucleotides and the DNA-binding protein  
 CC for a time sufficient to permit binding of the test molecule to test  
 CC sequences in the duplex DNA; (iii) separating unbound test  
 CC oligonucleotides from test oligonucleotides bound to binding protein;  
 CC (iv) amplifying the unbound test oligonucleotides; (v) repeating steps  
 CC (ii) to (iv); (vi) isolating the amplified test oligonucleotides; and  
 CC (vii) sequencing the isolated test oligonucleotides. Test sequences  
 CC AAX17001-X17481 and AAX17600 correspond to promoter targets for human  
 CC genes and test sequences AAX17482-X17599 correspond to promoter targets  
 CC for viral genes.  
 CC  
 SQ Sequence 50 BP; 18 A; 14 C; 10 G; 8 T; 0 other;

Query Match 88.9%; Score 16; DB 20; Length 50;  
 Best Local Similarity 100.0%; Pred. No. 5.7;  
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GGCTGCTGCACCTGT 16  
 Db 41 GGCTGCTGCACCTGT 26

RESULT 6  
 ABK83037/C  
 ID ABK83037 standard; DNA; 50 BP.  
 XX  
 AC ABK83037;  
 XX  
 DT 27-AUG-2002 (first entry)  
 XX  
 DE DNA binding molecule screening method test sequence #546.  
 XX  
 KM DNA binding molecule screening; inhibition of transcription;  
 KM infection; human immunodeficiency virus; HIV; parasite; cancer;  
 KM cardiovascular; respiratory; gastrointestinal; endocrine; metabolic;  
 KM rheumatic; immunological; haematological; neurological;  
 KM psychiatric; dermatological; ophthalmological; musculo-skeletal;  
 KM urogenital disorder; ss.  
 KM  
 OS Synthetic.  
 OS  
 PN US6384208-B1.  
 PN  
 PD 07-MAY-2002.  
 PD  
 PF 15-JUL-1999; 99US-0354947.  
 PF  
 XX 20-DEC-1993; 93US-0171389.  
 PR 07-JUN-1995; 95US-0482080.  
 PR 27-JUN-1991; 91US-0723618.  
 PR 23-DEC-1992; 92US-0996783.  
 PR 17-SEP-1993; 93US-0123936.  
 PR  
 XX  
 PA (GENE-) GENELABS TECHNOLOGIES INC.  
 PA  
 PI Edwards CA, Cantor CR, Andrews BM, Turin LM, Fry KE;  
 PI  
 XX WPI; 2002-442819/47.  
 DR  
 XX  
 PT Decreasing transcriptional activity of genes for treating infections or  
 PT cancer, by administration of an agent that binds to two non-overlapping  
 PT regions of the gene -  
 PT  
 XX  
 PS Example 15; SEQ ID No 546; 98pp; English.  
 PS  
 XX  
 CC The invention relates to a method of decreasing transcriptional activity  
 CC in a duplex deoxyribonucleic acid (DNA) template (T1) comprising  
 CC contacting (T1) with a binding agent comprising at least one small duplex  
 CC DNA-binding molecule (T2) coupled to at least one other small duplex-  
 CC binding molecule that binds to a non-overlapping region of target  
 CC sequence (T3). The method is useful for inhibiting transcription of a



CC range of disease-related genes for treating infections (by viruses,  
 CC including human immunodeficiency virus, bacteria, fungi, protozoa  
 CC and parasites), cancer, cardiovascular, respiratory, gastrointestinal,  
 CC endocrine/metabolic, rheumatic/immunological, hematological,  
 CC neurological, psychiatric, dermatological, ophthalmological,  
 CC musculo-skeletal, genetic or urogenital disorders. The method provides  
 CC sequence-specific inhibition of transcription of pathological genes  
 CC without affecting transcription of cellular genes regulated by the same  
 CC transcription factor, and can be applied to regulation of any gene.  
 CC ABK82492-ABK83155 represent DNA binding molecule test sequences used in  
 CC the method of the invention.

XX SQ Sequence 50 BP; 18 A; 14 C; 10 G; 8 T; 0 other;

Query Match 88.9%; Score 16; DB 24; Length 50;  
 Best Local Similarity 100.0%; Pred.No.5.7;  
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 GGCTGGTGCACCTGT 16  
 |||||  
 41 GGCTGGTGCACCTGT 26

RESULT 7  
 AAC01776/c

ID AAC01776 standard; cDNA; 435 BP.

XX AC AAC01776;

XX DT 06-OCT-2000 (first entry)

XX DE Human secreted protein 5' EST, SEQ ID NO: 1774.

XX KM Human; 5' EST; expressed sequence tag; secreted protein; cDNA isolation;  
 XX gene therapy; chromosome mapping; ss.

XX OS Homo sapiens.

XX PN EP1033401-A2.

XX PD 06-SEP-2000.

XX PF 21-FEB-2000; 2000EP-0200610.

XX PR 26-FEB-1999; 99US-0122487.

XX PA (GEST) GENSET.

XX PI Dumas Milne Edwards J, Duclert A, Giordano J;

XX PS WPI; 2000-500381/45.  
 P-PSDB; AAG01770.

XX PT New nucleic acid that is a 5' expressed sequence tag (5' EST) for  
 PT obtaining cDNAs and genomic DNAs that correspond to 5'ESTs and for  
 PT diagnostic, forensic, gene therapy and chromosome mapping procedures -  
 XX Claim 1; SEQ ID 1774; 71pp + CD-ROM; English.

XX The present sequence is one of a large number of 5' ESTs derived from  
 CC mRNA encoding secreted proteins. An ORF has been identified within the  
 CC sequence. The 5' ESTs were prepared from total human RNAs or polyA+ RNAs  
 CC derived from 30 different tissues. EST sequences usually correspond  
 CC mainly to the 3' untranslated region (UTR) of the mRNA because they are  
 CC often obtained from oligo-dT primed cDNA libraries. Such ESTs are not  
 CC well suited for isolating cDNA sequences derived from the 5' ends of  
 CC mRNAs and even in those cases where longer cDNA sequences have been  
 CC obtained, the full 5' UTR is rarely included. 5' ESTs are derived from  
 CC mRNAs with intact 5' ends and can therefore be used to obtain full length  
 CC cDNAs and genomic DNAs. 5' ESTs are also used in diagnostic, forensic,  
 CC gene therapy and chromosome mapping procedures. They are used to obtain  
 CC upstream regulatory sequences and to design expression and secretion  
 CC vectors.

XX SQ Sequence 435 BP; 105 A; 114 C; 133 G; 79 T; 4 other;

Query Match 88.9%; Score 16; DB 21; Length 435;  
 Best Local Similarity 100.0%; Pred.No.5.6;  
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 GGCTGGTGCACCTGT 16  
 |||||  
 DB 186 GGCTGGTGCACCTGT 171

RESULT 8  
 AAS61522/c

ID AAS61522 standard; cDNA; 455 BP.

XX AC AAS61522;

XX DT 29-JAN-2002 (first entry)

XX DE Lung small cell carcinoma antigen, cDNA #63.

XX KM Human; cytostatic; antitumour; lung small cell cancer antigen;  
 XX tumour; lung cancer; ss.

XX OS Homo sapiens.

XX PN WO200177168-A2.

XX PD 18-OCT-2001.

XX PF 11-APR-2001; 2001WO-US11859.

XX PR 11-APR-2000; 2000US-196780P.

XX PR 21-JUN-2000; 2000US-21381P.

XX PR 01-SEP-2000; 2000US-229763P.

XX PR 05-SEP-2000; 2000US-230629P.

XX PR 14-SEP-2000; 2000US-232565P.

XX PR 19-DEC-2000; 2000US-257037P.

XX PR 08-JUN-2001; 2001US-260796P.

XX PA (CORI-) CORIXA CORP.

XX PI Lodes MJ, Wang T, Mohamath R, Indirias CY;

XX PR WPI; 2002-010896/01.

XX DT Lung tumour polynucleotide and polypeptides useful in therapy and  
 XX diagnosis of cancer especially lung cancer -

XX Claim 1; Page 146; 295pp; English.

XX The invention relates to novel isolated lung small cell cancer antigen  
 CC polynucleotides (I) and polypeptides (II) used in a method of detecting  
 CC cancer in a patient. The method is optionally performed by  
 CC utilising oligonucleotides (III), where the biological sample  
 CC from the patient is contacted with (III), detecting the amount of  
 CC polynucleotide hybridized to (III) in the sample and comparing the  
 CC amount of polynucleotide to a predetermined cut-off value and thereby  
 CC determining cancer in a patient. (I), (II) or antigen-presenting cells  
 CC expressing (II) is useful for stimulating and/or expanding T cells  
 CC specific for a tumour protein. The method comprises contacting T cells  
 CC with one of the components under conditions to permit the stimulation  
 CC and/or expansion of the cells. A composition comprising (I) is useful for  
 CC stimulating an immune response in a patient and for inhibiting the  
 CC development of a cancer especially lung cancer in a patient. An  
 CC isolated T cell population is useful for removing tumour cells from the  
 CC biological sample and for inhibiting the development of cancer in a  
 CC patient. AAS6160-AAS61874 represent novel human lung small cell  
 CC cancer antigen coding sequences of the invention.

XX SQ Sequence 455 BP; 122 A; 116 C; 138 G; 78 T; 1 other;

Query Match 88.9%; Score 16; DB 24; Length 455;  
 Best Local Similarity 100.0%; Pred. No. 5.6;  
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGCTGTCACCTGT 16  
 DB 177 GGCTGTCACCTGT 162

## RESULT 9

AA189104  
 ID AA189104 standard; cDNA; 675 BP.

XX AA189104;

DT 06-NOV-2001 (first entry)

DE Human polynucleotide SEQ ID NO 9164.

XX Human; cytokine; cell proliferation; cell differentiation; gene therapy;

KW vaccine; peptide therapy; stem cell growth factor; haematopoiesis;

KM tissue growth factor; immunomodulatory; cancer; leukaemia;

XX nervous system disorders; arthritis; inflammation; ss.

OS Homo sapiens.

PN WO200164835-A2.

PD 07-SEP-2001.

PF 26-FEB-2001; 2001MO-US04927.

XX 28-FEB-2000; 2000US-0515126.

PR 18-MAY-2000; 2000US-0577409.

XX (HYSE-) HYSEQ INC.

PI Tang YT, Liu C, Drmanac RT;

XX WPI; 2001-514838/56.

DR P-PSDB; AA009173.

XX Isolated nucleic acid and polypeptides, useful for preventing

PT diagnosing and treating e.g. leukaemia, inflammation and immune

PT disorders -

XX Claim 1; SEQ ID NO 9164; 1399bp + Sequence Listing; English.

XX The invention relates to human polynucleotides (AA179941-AA193841) and

CC the encoded proteins (AA000010-AA013910) that exhibit activity relating to

CC cytokine, cell proliferation or cell differentiation or which may induce

CC production of other cytokines in other cell populations. The

CC polynucleotides and polypeptides are useful in gene therapy, vaccines or

CC peptide therapy. The polypeptides have various cytokine-like activities,

CC e.g. stem cell growth factor activity, haematopoiesis regulating

CC activity, tissue growth factor activity, immunomodulatory activity and

CC activin/inhibin activity and may be useful in the diagnosis and/or

CC treatment of cancer, leukaemia, nervous system disorders, arthritis and

CC inflammation.

CC Note: The sequence data for this patent did not form part of the printed

CC specification, but was obtained in electronic format directly from WIPO

CC at ftp.wipo.int/pub/published\_pct\_sequences.

XX Sequence 675 BP; 189 A; 165 C; 147 G; 172 T; 2 other;

QY Query Match 88.9%; Score 16; DB 22; Length 675;

Best Local Similarity 100.0%; Pred. No. 5.5;

Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

DB 3 CTGATGTCACCTGTTA 18

437 CTGATGTCACCTGTTA 452

RESULT 10  
 ID AA43611/C  
 AA43611 standard; DNA; 1142 BP.

XX AA43611;

DT 24-SEP-1998 (first entry)

DE Human secreted protein 11 encoding DNA.

XX Secreted protein; human; cell proliferation; cytokine activity;

KW tissue growth; cellular differentiation; regeneration; activin;

KM inhibin; chemotactic; haemostatic; thrombolytic; tumour inhibition;

XX anti-inflammatory activity; biomarker; ss.

OS Homo sapiens.

PN Key Location/Qualifiers

XX CDS 33..788

XX /tag= a

XX /product= "human secreted protein"

XX 11-DEC-1997; 97MO-US22787.

XX 18-JUN-1998.

XX 11-DEC-1996; 96US-0032757.

XX (CHIR ) CHIRON CORP.

XX Escobedo J, Garcia P, Hu Q, Kothakota S, Williams LT;

XX WPI; 1998-348453/30.

DR P-PSDB; AA63691.

XX Secreted human polypeptides - having cytokine, cell proliferation or

PT differentiation, activin or inhibin, tumour inhibition or

PT anti-inflammatory activities

XX Claim 6; Page 38; 78bp; English.

XX This DNA encodes a human secreted protein. The specification provides

CC secreted protein sequences (AA63681 to AA63699) encoded by the nucleic

CC acid sequences shown in AA43611 to AA43619. The invention provides a

CC method of identifying a secreted polypeptide which is modified by rough

CC microsomes. The secreted proteins can be used in assays to determine

CC biological activities, such as cytokine, cell proliferation, or cellular

CC differentiation activities, tissue growth or regeneration, activin or

CC inhibin activity, chemotactic or chemokinetic activity, haemostatic or

CC thrombolytic activity, receptor/ligand activity, tumour inhibition, or

CC anti-inflammatory activity. The proteins can also be used as

CC biomarkers, to identify tissues or cell types which express the proteins,

CC or a stage- or disease-specific alteration in protein expression. They

CC can be used in protein interaction assays, to identify ligands or binding

CC proteins. Compounds which affect the biological activities of the

CC secreted proteins or their ability to interact with specific ligands can

CC be identified using the proteins in screening assays. The proteins and

CC antibodies that bind specifically to the protein can also be used to

CC design diagnostic tests and therapeutic compositions for diseases which

CC may be associated with altered expression of these proteins. Fusion

CC proteins comprising, e.g. signal sequences or transmembrane domains of

CC the proteins can be used to target other protein domains to cellular

CC membrane or they can be secreted extracellularly.

XX Sequence 1142 BP; 304 A; 335 C; 331 G; 172 T; 0 other;

QY Query Match 88.9%; Score 16; DB 19; Length 1142;

Best Local Similarity 100.0%; Pred. No. 5.5;

Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGCTGTCACCTGT 16  
 |||||  
 DB 159 GGCTGTCACCTGT 144

RESULT 11  
 ID AAV62754/c  
 AAV62754 standard; cDNA, 1152 BP.

XX AAV62754;

DT 15-FEB-1999 (first entry)

XX Human secreted protein clone fml50\_1 cDNA.

XX Secreted protein; human; fml50\_1; ds.

XX Homo sapiens.

XX Key Location/Qualifiers

FT CDS 11..973

FT /tag= a

XX WO9846757-A2.

XX 22-OCT-1998.

XX 14-APR-1998; 98MO-US07999.

XX 13-APR-1998; 98US-0059487.

XX 15-APR-1997; 97US-0843374.

XX (GENY ) GENETICS INST INC.

XX Agostino MJ, Jacobs K, Lavallie ER, McCoy JM, Merberg D,

XX Racie LA, Spaulding V, Treacy M,

XX P-PSDB; AAM74728.

XX WPI; 1998-568731/48.

XX Claim 34; Page 95; 120pp; English.

XX Full-length cDNA clone fml50\_1 includes an open reading frame

XX encoding a human secreted protein (see AAM74728). It was isolated

XX from a human adult brain cDNA library using methods which are

XX selective for cDNAs encoding secreted proteins, or was identified

XX as encoding a secreted or transmembrane protein on the basis of

XX computer analysis of the amino acid sequence of the encoded protein.

XX It shows some similarity to database sequences. The invention

XX provides polynucleotides (see AAV62746-55) from human foetal brain,

XX adult testis, adult brain, adult kidney and foetal kidney (all

XX deposited as composite clone ATCC 98404), which encode human

XX secreted proteins (see AAM74720-29). The polynucleotides and

XX make them suitable for treating, preventing or ameliorating medical

XX conditions in humans and animals, although no supporting data are

XX given. Suggested activities include nutritional activity, immune

XX stimulating (e.g. as vaccines) or suppressing activity,

XX haemotopoiesis regulating activity, tissue growth activity,

XX activin/inhibin activity, chemotactic/chemokinetic activity,

XX haemostatic and thrombolytic activity, receptor/ligand activity,

XX antiinflammatory activity, cadherin/tumour invasion suppressor

XX activity, and tumour inhibition activity. The polynucleotides are

XX also stated to be useful for gene therapy, and can be used in

XX recombinant production of the polypeptides.

XX Sequence 1152 BP; 342 A; 329 C; 317 G; 164 T; 0 other;

XX Query Match 88.9%; Score 16; DB 19; Length 1152;

Best Local Similarity 100.0%; Pred. No. 5.5;  
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGCTGTCACCTGT 16  
 |||||  
 DB 137 GGCTGTCACCTGT 122

RESULT 12  
 ID ABQ92057/c  
 ABQ92057 standard; cDNA, 1152 BP.

XX ABQ92057;

DT 04-OCT-2002 (first entry)

XX Human polynucleotide SEQ ID NO 54.

XX Human; cytosolic; antirheumatic; antiarthritic; vulnery; analgesic;

XX antiinflammatory; antibacterial; immunosuppressive; antiparkinsonian;

XX neuroprotective; nootropic; osteopathic; haemostatic; vasotropic;

XX antitumor; fungicide; antidiabetic; antiaesthetic; antiallergic;

XX immunostimulant; antiparasitic; secreted protein; transmembrane protein;

XX cytokine; cell proliferation; cell differentiation; autoimmune disease;

XX stem cell; growth factor; nervous system disease; neuropathy;

XX Alzheimer's disease; Parkinson's disease; Huntington's disease;

XX osteoporosis; severe combined immunodeficiency; SCID; infection;

XX multiple sclerosis; rheumatoid arthritis; gene therapy; gene; ss.

XX Homo sapiens.

XX US2002065394-A1.

XX 30-MAY-2002.

XX 22-DEC-2000; 2000US-0745763.

XX 18-MAR-1998; 98US-0040963.

XX (JACO/) JACOBS K.

XX (MCCO/) MCCOY J M.

XX (LAVA/) LAVALLIE E R.

XX (COLL/) COLLINS-RACIE L A.

XX (EVAN/) EVANS C.

XX (MERB/) MERBERG D.

XX (TREAC/) TREACY M.

XX (SPAUL/) SPAULDING V.

XX Jacobs K, McCoy JM, Lavallie ER, Collins-Racie LA, Evans C,

XX Merberg D, Treacy M, Spaulding V;

XX WPI; 2002-562343/62.

XX P-PSDB; ABP61843.

XX Novel secreted or transmembrane protein and polynucleotide encoding the

XX protein, useful for diagnosis and treatment of neurological disorders,

XX cancer, autoimmune diseases, bone disorders and lung or liver fibrosis

XX Claim 189; Page 195-196; 284pp; English.

XX The invention relates to human secreted or transmembrane protein (I),

XX their fragments and is encoded by specific complementary deoxyribonucleic

XX acid (cDNA) inserts (II), where the protein is substantially free from

XX other mammalian proteins. (I) are useful for preventing, treating or

XX ameliorating a medical condition, especially immunological treatment or

XX prevention of tumors. (I) exhibits activity relating to angiogenesis,

XX cytokine, cell proliferation, cell differentiation, antiinflammatory,

XX stem cell growth factor activity and activin or inhibin-related

XX activities. (I) can be used to manipulate stem cells in culture to give

XX rise to neuroepithelial cells that can be used to augment or replace

XX cells damaged by illness, autoimmune disease, accidental damage or

XX genetic disorders. (I) induces the proliferation of neural cells and

regeneration of nerve and brain tissue and is useful for the treatment of central and peripheral nervous system diseases and neuropathies, such as Alzheimer's, Parkinson's disease, Huntington's disease, amyotrophic lateral sclerosis. (I) is involved in chemotactic or chemokinetic activity, regulation of haematopoiesis and is useful for treating myeloid CC or lymphoid cell disorders, platelet disorders such as thrombocytopenia CC and for regeneration of bone, cartilage, tendon, ligament and/or nerve CC tissue growth and in tissue repair, healing of burns, incisions, ulcers, CC for treating osteoporosis, osteoarthritis, bone degenerative disorders or CC periodontal disease. (II) is also useful for gut protection or CC regeneration and treatment of lung or liver fibrosis, reperfusion injury CC in various tissues, various immune deficiencies and disorders including CC severe combined immunodeficiency (SCID), bacterial or fungal infections, CC autoimmune disorders e.g. multiple sclerosis, rheumatoid arthritis, CC diabetes mellitus, myasthenia gravis, allergic reactions and conditions, CC such as asthma or other respiratory problems. (II) is useful to express CC recombinant protein, as markers for tissues in which the corresponding CC protein is preferentially expressed and in gene therapy. The present CC sequence is that of a polynucleotide of the invention.

SO Sequence 1152 BP; 342 A; 329 C; 317 G; 164 T; 0 other;

Query Match 88.9%; Score 16; DB 24; Length 1152;  
Best Local Similarity 100.0%; Pred. No. 5.5;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 GGCTGGTGCACCTGT 16  
|||  
Db 137 GGCTGGTGCACCTGT 122

RESULT 13  
AAC77733/c  
ID AAC77733 standard; cDNA; 1217 BP.

AC AAC77733;

DT 08-FEB-2001 (first entry)

DE Human cancer associated gene sequence SEQ ID NO:127.

XX Human; cancer associated gene; cancer antigen; detection; cancer;  
XX diagnosis; cytostatic; proliferative; vulnerrary; immunomodulator;  
XX antiadhesive; antidiabetic; antirheumatic; antiarthritic; antiviral;  
XX antiinflammatory; antithyroid; antiallergic; antibacterial; cardiac;  
XX dermatological; neuroprotective; thrombolytic; coagulant; nocropic;  
XX vasotropic; antipsoriatic; antiangiogenic; gene therapy; inflammation;  
XX immune disorder; hematopoietic cell disorder; autoimmune disorder;  
XX allergic reaction; graft versus host disease; organ rejection;  
XX haemostatic; thrombolytic; cardiovascular disorder; infection;  
XX neurological disease; drug screening; ss.

OS Homo sapiens.

XX MO200055350-A1.

2

XX 21-SEP-2000.

XX 08-MAR-2000; 2000MO-US05882.

XX 12-MAR-1999; 99US-0124270.

XX (HUMA-) HUMAN GENOME SCI INC.

XX Rosen CA, Ruben SM;

XX WPI; 2000-587533/55.

XX P-PSDB; AAB3524.

XX Novel isolated nucleic acids comprising sequences encoding peptides  
XX useful for treating or diagnosing e.g. cancer -

XX Claim 1; Page 709; 2352pp; English.

XX AAC77607 to AAC78448 encode the human cancer associated proteins given  
CC in AAB43398 to AAB44239. The proteins can have activities based on the  
CC tissues and cells the genes are expressed in. Example of activities  
CC include: cytostatic; proliferative; vulnerrary; immunomodulator;  
CC antidiabetic; antiasclmatic; antirheumatic; antiarthritic;  
CC antiinflammatory; antithyroid; antiallergic; antibacterial; antiviral;  
CC dermatological; neuroprotective; cardiac; thrombolytic; coagulant;  
CC nocropic; vasotropic; antipsoriatic and antiangiogenic. The  
CC polynucleotides and polypeptides can be used for preventing, treating or  
CC ameliorating medical conditions and diagnosing pathological conditions.  
CC Polynucleotides, polypeptides, antibodies, agonists and antagonists from  
CC the present invention may be used to treat immune disorders by activating  
CC or inhibiting the proliferation, differentiation or mobilisation of  
CC immune cells, to treat disorders of haematopoietic cells, autoimmune  
CC disorders, allergic reactions, graft versus host disease and organ  
CC rejection, modulate haemostatic or thrombolytic activity, modulate  
CC inflammation, cancers, cardiovascular disorders, neurological disease and  
CC bacterial or viral infections. The peptides, nucleotides, antibodies, and  
CC agonists and antagonists may be also be used in drug screens. AAC78449 to  
CC AAC78457 and AAB44240 represent sequences used in the exemplification of  
CC the present invention.

SO Sequence 1217 BP; 300 A; 360 C; 367 G; 187 T; 3 other;

Query Match 88.9%; Score 16; DB 21; Length 1217;  
Best Local Similarity 100.0%; Pred. No. 5.5;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 GGCTGGTGCACCTGT 16  
|||  
Db 235 GGCTGGTGCACCTGT 220

RESULT 14  
AAZ17770  
ID AAZ17770 standard; cDNA; 1278 BP.

AC AAZ17770;

DT 12-OCT-1999 (first entry)

DE Human gene expression product cDNA sequence SEQ ID NO:5243.

XX Human; gene; gene expression product; diagnosis; therapy; probe;  
XX detection; mapping; tissue typing; profiling; forensic; cancer;  
XX genetic analysis; colorectal cancer; breast cancer; lung cancer; ss.

XX Homo sapiens.

XX MO9938972-A2.

XX 05-AUG-1999.

XX 28-JAN-1999; 99WO-US01619.

XX 03-APR-1998; 98US-0080666.

XX 28-JAN-1998; 98US-0072910.

XX 24-FEB-1998; 98US-0075954.

XX 31-MAR-1998; 98US-0080114.

XX 03-APR-1998; 98US-0080515.

XX (CHIR ) CHIRON CORP.

XX (HYSE-) HYSEQ INC.

XX Ctkvenjakov R, Dickson M, Drmanac R, Drmanac S;  
XX Escobedo J, Garcia PD, Garcia V, Glese K, Innis MA;  
XX Jones WL, Kassam A, Kennedy GC, Kita D, Labat I;  
XX Lamson G, Leshkowitz D, Pot D, Randazzo F, Reinhard C;  
XX Stache-Crahn B, Suduth-Klinger J, Williams LR;  
XX WPI; 1999-494092/41.

PT Novel human genes and their expression products which are  
PT differentially expressed in different cell types  
PS Claim 1: Page 2475-2476; 2479pp; English.

CC The present invention describes a library of human polynucleotides  
CC comprising the sequences given in AA212532 to AA217779. Also described is  
CC a method of detecting differentially expressed genes correlated with the  
CC cancerous state of a mammalian cell, comprising detecting at least one  
CC differentially expressed gene product in a test sample from a cell  
CC suspected of being cancerous, where the gene product is encoded by one  
CC of the 5248 polynucleotide sequences given in AA212532 to AA217779. The  
CC polynucleotides can be used as a source of primers and probes, which can  
CC be used for a variety of purposes, e.g. detection of expression levels,  
CC mapping, tissue typing or profiling, forensics, genetic analysis and  
CC detection of polymorphisms. Polypeptides encoded by the polynucleotides  
CC can be used for raising antibodies for experimental, diagnostic and  
CC therapeutic purposes. The polynucleotides may also be used to construct  
CC arrays for diagnostics (which may be used to determine function of an  
CC encoded protein); and to detect differences in expression levels between  
CC two cells (e.g. to identify abnormal or diseased tissue in a human, to  
CC identify a genetic predisposition or susceptibility to a disease such as  
CC cancer). The polynucleotides of the invention are especially used in the  
CC diagnosis, prognosis and management of colorectal cancer, breast cancer,  
CC and lung cancer. The polynucleotides can also be used to screen for  
CC peptide analogues and antagonists.

XX Sequence 1278 BP; 216 A; 383 C; 387 G; 282 T; 10 other;

Query Match 88.9%; Score 16; DB 20; Length 1278;

Best Local Similarity 100.0%; Pred. No. 5.5;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGCTGGTGCACCTGT 16  
DB 1118 GGCTGGTGCACCTGT 1133

RESULT 15

ACC46555/c

ID ACC46555 standard; cDNA; 2382 BP.

AC ACC46555;

XX 02-JUN-2003 (first entry)

XX Human dithp secreted/extracellular matrix protein-encoding cDNA.

KW Human; dithp; diagnostic and therapeutic polynucleotide; diagnosis;  
KW cancer; cell proliferative disorder; autoimmune disorder;  
KW inflammatory disorder; infection; hormonal disorder; metabolic disorder;  
KW neurological disorder; gastrointestinal disorder; transport disorder;  
KW connective tissue disorder; drug screening; proteome analysis;  
KW gene therapy; antisense therapy; genotyping; transgenic animal; knock in;  
KW disease model; toxicological testing; transcript imaging;  
KW secreted protein; extracellular matrix; gene; ss.

XX Homo sapiens.

XX WO200297031-A2.

XX 05-DEC-2002.

XX 27-MAR-2002; 2002MO-US10056.

XX 28-MAR-2001; 2001US-279619P.

XX 29-MAR-2001; 2001US-280067P.

XX 29-MAR-2001; 2001US-280068P.

XX 16-MAY-2001; 2001US-281280P.

XX 17-MAY-2001; 2001US-281829P.

XX 17-MAY-2001; 2001US-291849P.

XX 19-JUN-2001; 2001US-299428P.

XX 20-JUN-2001; 2001US-299776P.

PR 20-JUN-2001; 2001US-300001P.  
XX (INCY-) INCYTE GENOMICS INC.

PI Dafio A, Jones AL, Tran AB, Dahl CR, Gietzen D, Chinn J;  
PI DuFour GE, Hillman JL, Yu JY, Tuason O, Yap PE, Anshay SR;  
PI Daugterey SC, Dan TC, Liu TF, Nguyen DA, Kleefeld Y, Gertlin EH,  
PI Peralta CH, David MH, Lewis SA, Chen AJ, Panzer SR, Harris B,  
PI Flores V, Marwaha R, Lo A, Lan RV, Urashka ME;  
XX WPI; 2003-129518/12.  
DR P-PsDB; ABR41618.

PT Novel human diagnostic and therapeutic polypeptide useful for  
PT identifying test compound which specifically binds to a polypeptide  
PT encoded by human diagnostic and therapeutic polynucleotide, and to  
PT induce antibodies

XX Claim 2; SEQ ID No 476; 591pp; English.

CC The invention relates to novel human diagnostic and therapeutic  
CC polynucleotides designated dithp (ACC46080-ACC46749) and to their  
CC encoded proteins (DITHP; ABR41136-ABR41812). The invention also relates  
CC to polynucleotide sequences at least 90% identical to the dithp cDNA  
CC sequences of the invention; recombinant vectors, host cells and  
CC transgenic organisms comprising a dithp nucleic acid sequence; the  
CC recombinant production of DITHP proteins; antibodies specific for DITHP  
CC proteins; microarrays comprising dithp nucleic acid sequences; methods  
CC for detecting dithp nucleotide and protein sequences; methods of screening  
CC for compounds which specifically bind a DITHP protein; and methods of  
CC assessing the toxicity of test compounds using a dithp hybridisation  
CC probe. Dithp nucleic acid sequences and DITHP proteins may be used in the  
CC diagnosis of a wide variety of conditions including cancer and other cell  
CC proliferative disorders; autoimmune or inflammatory disorders; bacterial,  
CC viral, fungal or parasitic infections; hormonal disorders; metabolic  
CC disorders; neurological disorders; gastrointestinal disorders; transport  
CC disorders; and connective tissue disorders. They may also be used to  
CC screen for modulators of protein activity or gene expression. DITHP  
CC proteins can additionally be used in analysis of the proteome of a tissue  
CC or cell type and to induce antibodies. The dithp nucleic acids are  
CC additionally useful in somatic or germline gene therapy of the disorders  
CC mentioned above, as a source of antisense sequences, as a source of  
CC probes and primers, in genotyping and identification of individuals, in  
CC the generation of transgenic animal models of human disease or knock in  
CC humanised animals, in toxicological testing, and in transcript imaging.  
CC The present sequence represents a dithp cDNA encoding a DITHP protein.  
CC Note: The sequence data for this patent did not form part of the printed  
CC specification, but was obtained in electronic format directly from WIPO  
CC at ftp.wipo.int/pub/published\_pct\_sequences.

XX Sequence 2382 BP; 456 A; 1032 C; 337 G; 556 T; 1 other;

Query Match 88.9%; Score 16; DB 25; Length 2382;

Best Local Similarity 100.0%; Pred. No. 5.4;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGCTGGTGCACCTGT 16  
DB 1869 GGCTGGTGCACCTGT 1854

RESULT 16

AAAN50114/c

ID AAAN50114 standard; DNA; 2721 BP.

XX AAAN50114;

XX 25-MAR-2003 (updated)

XX 17-OCT-1991 (first entry)

XX DNA sequence encoding Epstein-Barr virus (EBV) outer surface protein.

```
KM Epstein-Barr virus; antigen; vaccine; ss.
XX
XX Epstein-Barr virus.
XX
XX Key Location/Qualifiers
FT mat_peptide 1..2721
FT /*tag= a
FT /label= EBV surface protein antigen
XX
XX EP151079-A.
XX
XX 07-AUG-1985.
XX
XX 28-JAN-1985; 85EP-0400141.
XX
XX 23-JUL-1984; 84US-0633558.
XX 30-JAN-1984; 84US-0575352.
XX
XX (UYCH-) UNIV CHICAGO.
XX
XX Kieff E, Tanner J, Hummel M, Belset C;
XX
XX WPI; 1985-191978/32.
XX P-PSDB; AAP50073.
XX
XX New fragment of Epstein-Barr Virus DNA - useful in vector to
XX express polypeptide for use in prepn. of vaccine against the
XX virus and for use in diagnosis.
XX
XX Claim 1; Page 21-23; 26pp; English.
XX
XX The sequence encodes an outer surface viral protein of EBV, used
XX to generate antibodies reacting with the surface proteins of
XX EBV-infected cells, and in the preparation of a vaccine against EBV.
XX (Updated on 25-MAR-2003 to correct PA field.)
XX
XX Sequence 2721 BP; 762 A; 876 C; 557 G; 526 T; 0 other;
SQ
Query Match 88.9%; Score 16; DB 6; Length 2721;
Best Local Similarity 100.0%; Pred. No. 5.4;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 GGCTGGTGCACCTGT 16
DB 2124 GGCTGGTGCACCTGT 2109
RESULT 17
AAT04821/c
ID AAT04821 standard; cDNA; 5931 BP.
XX
XX AAT04821;
AC
XX 18-JAN-1996 (first entry)
XX
XX EBV gp350/220 cDNA.
DE
XX EBV, gp350; gp220; gp350/gp220; non-splicing variant; vaccine; ds.
XX
XX Epstein-Barr virus.
XX
XX Key Location/Qualifiers
FT CDS 1014..3737
FT /*tag= a
FT sig_peptide 1014..1067
FT /*tag= b
FT mat_peptide 1068..3734
FT /*tag= c
FT misc_feature 2514..2515
FT /*tag= d
FT /function= splice donor site
FT /note= "Bases 2513-2517 (AAGT) are replaced by
FT GTCA in the non-splicing variant"
```

```
FT misc_feature 3105..3106
FT /*tag= e
FT /function= splice acceptor site
FT /note= "Bases 3104-3107 (AGGT) are replaced by
FT polyA_signal 3742..3747
FT /*tag= f
XX
XX MO9528488-A1.
XX
XX 26-OCT-1995.
XX
XX 13-APR-1995; 95WO-US04611.
XX
XX 18-APR-1994; 94US-0229291.
XX
XX (AVIR-) AVIRON.
XX
XX Jackman WT, Spaete R;
XX
XX WPI; 1995-373802/48.
XX P-PSDB; AAR80144.
XX
XX New DNA encoding a homogeneous gp350 protein - can be used for
XX preventing and treating Epstein-Barr virus-related diseases or
XX conditions
XX
XX Claim 2; Fig.1; 61pp; English.
XX
XX The donor and acceptor splice sites of the EBV gene encoding gp350/
XX 220 are mutated by replacement of native nucleotides by non-native
XX nucleotides, without altering the encoded amino acid sequence.
XX resulting in elimination of gp220 prodn. Recombinant homogeneous
XX gp350, useful in vaccines, is expressed in mammalian or insect cell
XX hosts.
XX
XX Sequence 5931 BP; 1453 A; 1782 C; 1437 G; 1259 T; 0 other;
SQ
Query Match 88.9%; Score 16; DB 16; Length 5931;
Best Local Similarity 100.0%; Pred. No. 5.4;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 GGCTGGTGCACCTGT 16
DB 3137 GGCTGGTGCACCTGT 3122
RESULT 18
ABA15144
ID ABA15144 standard; DNA; 13559 BP.
XX
XX ABA15144;
AC
XX 23-JAN-2002 (first entry)
XX
XX Human nervous system related polynucleotide SEQ ID NO 7475.
DE
XX
XX Human; nocotropic; neuroprotective; cytosaric; dermatological; virucide;
XX immunosuppressive; antiinflammatory; anti-HIV; antibacterial; vulnerary;
XX antiparkinsonian; antischling; antianaemic; antiarthritic; cancer;
XX antirheumatic; hepatotropic; cerebroprotective; antiinflammatory;
XX antiallergic; antidiabetic; antidiuretic; anticonvulsant; antifungal;
XX antiparasitic; cardiant; immune disorder; cardiovascular disorder;
XX neurological disease; infection; nephrotropic; gene therapy; vaccine; ds.
XX
XX Homo sapiens.
XX
XX MO200159063-A2.
XX
XX 16-AUG-2001.
XX
XX 17-JAN-2001; 2001WO-US01334.
XX
```

PR 31-JAN-2000; 2000US-0179065.  
 PR 04-FEB-2000; 2000US-0180628.  
 PR 24-FEB-2000; 2000US-0484664.  
 PR 02-MAR-2000; 2000US-0186350.  
 PR 16-MAR-2000; 2000US-0189874.  
 PR 17-MAR-2000; 2000US-0190076.  
 PR 18-APR-2000; 2000US-0198123.  
 PR 19-MAY-2000; 2000US-0205515.  
 PR 07-JUN-2000; 2000US-0209467.  
 PR 28-JUN-2000; 2000US-0214886.  
 PR 30-JUN-2000; 2000US-0215135.  
 PR 07-JUL-2000; 2000US-0216647.  
 PR 07-JUL-2000; 2000US-0216880.  
 PR 11-JUL-2000; 2000US-0217487.  
 PR 11-JUL-2000; 2000US-0217496.  
 PR 14-JUL-2000; 2000US-0218290.  
 PR 26-JUL-2000; 2000US-0220963.  
 PR 26-JUL-2000; 2000US-0220964.  
 PR 14-AUG-2000; 2000US-0224518.  
 PR 14-AUG-2000; 2000US-0224519.  
 PR 14-AUG-2000; 2000US-0225213.  
 PR 14-AUG-2000; 2000US-0225214.  
 PR 14-AUG-2000; 2000US-0225266.  
 PR 14-AUG-2000; 2000US-0225267.  
 PR 14-AUG-2000; 2000US-0225268.  
 PR 14-AUG-2000; 2000US-0225270.  
 PR 14-AUG-2000; 2000US-0225447.  
 PR 14-AUG-2000; 2000US-0225757.  
 PR 14-AUG-2000; 2000US-0225758.  
 PR 14-AUG-2000; 2000US-0225759.  
 PR 18-AUG-2000; 2000US-0226279.  
 PR 22-AUG-2000; 2000US-0226681.  
 PR 22-AUG-2000; 2000US-0226688.  
 PR 22-AUG-2000; 2000US-0227182.  
 PR 23-AUG-2000; 2000US-0227009.  
 PR 30-AUG-2000; 2000US-0228924.  
 PR 01-SEP-2000; 2000US-0229287.  
 PR 01-SEP-2000; 2000US-0229343.  
 PR 01-SEP-2000; 2000US-0229344.  
 PR 01-SEP-2000; 2000US-0229345.  
 PR 05-SEP-2000; 2000US-0229509.  
 PR 05-SEP-2000; 2000US-0229513.  
 PR 06-SEP-2000; 2000US-0230437.  
 PR 06-SEP-2000; 2000US-0230438.  
 PR 08-SEP-2000; 2000US-0231242.  
 PR 08-SEP-2000; 2000US-0231243.  
 PR 08-SEP-2000; 2000US-0231244.  
 PR 08-SEP-2000; 2000US-0231413.  
 PR 08-SEP-2000; 2000US-0231414.  
 PR 08-SEP-2000; 2000US-0232080.  
 PR 08-SEP-2000; 2000US-0232081.  
 PR 12-SEP-2000; 2000US-0231968.  
 PR 14-SEP-2000; 2000US-0232397.  
 PR 14-SEP-2000; 2000US-0232398.  
 PR 14-SEP-2000; 2000US-0232399.  
 PR 14-SEP-2000; 2000US-0232400.  
 PR 14-SEP-2000; 2000US-0232401.  
 PR 14-SEP-2000; 2000US-0233063.  
 PR 14-SEP-2000; 2000US-0233064.  
 PR 14-SEP-2000; 2000US-0233065.  
 PR 21-SEP-2000; 2000US-0234423.  
 PR 21-SEP-2000; 2000US-0234474.  
 PR 25-SEP-2000; 2000US-0234997.  
 PR 25-SEP-2000; 2000US-0234998.  
 PR 26-SEP-2000; 2000US-0235844.  
 PR 27-SEP-2000; 2000US-0235834.  
 PR 27-SEP-2000; 2000US-0235836.  
 PR 29-SEP-2000; 2000US-0236327.  
 PR 29-SEP-2000; 2000US-0236367.  
 PR 29-SEP-2000; 2000US-0236368.  
 PR 29-SEP-2000; 2000US-0236369.  
 PR 29-SEP-2000; 2000US-0236370.  
 PR 02-OCT-2000; 2000US-0236802.

PR 02-OCT-2000; 2000US-0237037.  
 PR 02-OCT-2000; 2000US-0237038.  
 PR 02-OCT-2000; 2000US-0237039.  
 PR 02-OCT-2000; 2000US-0237040.  
 PR 13-OCT-2000; 2000US-0239935.  
 PR 13-OCT-2000; 2000US-0239937.  
 PR 20-OCT-2000; 2000US-0240960.  
 PR 20-OCT-2000; 2000US-0241785.  
 PR 20-OCT-2000; 2000US-0241786.  
 PR 20-OCT-2000; 2000US-0241787.  
 PR 20-OCT-2000; 2000US-0241808.  
 PR 20-OCT-2000; 2000US-0241809.  
 PR 20-OCT-2000; 2000US-0241826.  
 PR 20-OCT-2000; 2000US-0242221.  
 PR 01-NOV-2000; 2000US-0244617.  
 PR 08-NOV-2000; 2000US-0246474.  
 PR 08-NOV-2000; 2000US-0246475.  
 PR 08-NOV-2000; 2000US-0246476.  
 PR 08-NOV-2000; 2000US-0246477.  
 PR 08-NOV-2000; 2000US-0246478.  
 PR 08-NOV-2000; 2000US-0246523.  
 PR 08-NOV-2000; 2000US-0246524.  
 PR 08-NOV-2000; 2000US-0246525.  
 PR 08-NOV-2000; 2000US-0246526.  
 PR 08-NOV-2000; 2000US-0246527.  
 PR 08-NOV-2000; 2000US-0246528.  
 PR 08-NOV-2000; 2000US-0246532.  
 PR 08-NOV-2000; 2000US-0246609.  
 PR 08-NOV-2000; 2000US-0246610.  
 PR 08-NOV-2000; 2000US-0246611.  
 PR 08-NOV-2000; 2000US-0246613.  
 PR 17-NOV-2000; 2000US-0249207.  
 PR 17-NOV-2000; 2000US-0249208.  
 PR 17-NOV-2000; 2000US-0249209.  
 PR 17-NOV-2000; 2000US-0249210.  
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 PR 17-NOV-2000; 2000US-0249212.  
 PR 17-NOV-2000; 2000US-0249213.  
 PR 17-NOV-2000; 2000US-0249214.  
 PR 17-NOV-2000; 2000US-0249215.  
 PR 17-NOV-2000; 2000US-0249216.  
 PR 17-NOV-2000; 2000US-0249217.  
 PR 17-NOV-2000; 2000US-0249218.  
 PR 17-NOV-2000; 2000US-0249244.  
 PR 17-NOV-2000; 2000US-0249245.  
 PR 17-NOV-2000; 2000US-0249264.  
 PR 17-NOV-2000; 2000US-0249265.  
 PR 17-NOV-2000; 2000US-0249297.  
 PR 17-NOV-2000; 2000US-0249299.  
 PR 17-NOV-2000; 2000US-0249300.  
 PR 01-DEC-2000; 2000US-0250391.  
 PR 01-DEC-2000; 2000US-0251160.  
 PR 05-DEC-2000; 2000US-0251030.  
 PR 05-DEC-2000; 2000US-0251988.  
 PR 05-DEC-2000; 2000US-0256719.  
 PR 06-DEC-2000; 2000US-0251479.  
 PR 08-DEC-2000; 2000US-0251856.  
 PR 08-DEC-2000; 2000US-0251868.  
 PR 08-DEC-2000; 2000US-0251869.  
 PR 08-DEC-2000; 2000US-0251989.  
 PR 08-DEC-2000; 2000US-0251990.  
 PR 11-DEC-2000; 2000US-0254097.  
 PR 05-JAN-2001; 2001US-0259678.  
 PA (HUMA-) HUMAN GENOME SCI INC.  
 XX  
 PI Rosen CA, Barash SC, Ruben SM;  
 XX WPI, 2001-541565/50.  
 DR Nucleic acids encoding 3224 human nervous system antigen polypeptides,  
 XX useful for preventing, diagnosing and/or treating nervous system  
 PT cancers and metastases -

XX Disclosure; SEQ ID NO 7475; 1701bp + Sequence Listing; English.  
PS  
XX  
CC The invention relates to novel genes (ABAI1004-ABA21534) and proteins  
CC (ABH14678-ABH18001) useful for preventing, treating or ameliorating  
CC medical conditions e.g. by protein or gene therapy. The genes are  
CC isolated from a range of human tissues disclosed in the specification.  
CC The nucleic acids, proteins, antibodies and (ant)agonists are useful  
CC in the diagnosis, treatment and prevention of: (a) cancer, e.g. breast  
CC and ovarian cancer and other cancers of the adrenal gland, bone, bone  
CC marrow, breast, gastrointestinal tract, liver, lung, or urogenital;  
CC (b) immune disorders e.g. Addison's disease, allergies, autoimmune  
CC hemolytic anaemia, autoimmune thyroiditis, diabetes mellitus, Crohn's  
CC colitis; (c) cardiovascular disorders such as myocardial ischaemia;  
CC (d) wound healing; (e) neurological diseases e.g. cerebral anoxia and  
CC epilepsy; and (f) infectious diseases such as viral, bacterial, fungal  
CC Note: The sequence data for this patent did not form part of the  
CC printed specification, but was obtained in electronic format directly  
CC from WIPO at [ftp.wipo.int/pub/published\\_pct\\_sequences](http://ftp.wipo.int/pub/published_pct_sequences).  
XX  
SQ Sequence 13559 BP; 2882 A; 3867 C; 4239 G; 2571 T; 0 other;  
Query Match 88.9%; Score 16; DB 22; Length 13559;  
Best Local Similarity 100.0%; Pred. No. 5.3;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
OY 1 GGCTGGTGTACCTGT 16  
Db 12761 GGCTGGTGTACCTGT 12776  
RESULT 19  
ABNS2794  
XX ABN52794 standard; DNA; 65 BP.  
XX  
AC ABN52794;  
XX  
DT 15-JUL-2002 (first entry)  
XX  
DE Mouse spliced transcript detection oligonucleotide SEQ ID NO:25542.  
XX  
XX Human: mouse; rat; splice transcript; detection; RNA transcript;  
XX splice variant; transcriptome; oligonucleotide library; ss.  
XX  
XX Mus musculus.  
XX  
XX OS  
XX  
XX PN WO200210449-A2.  
XX  
XX PD 07-FEB-2002.  
XX  
XX PF 20-JUL-2001; 2001WO-1B01903.  
XX  
XX PR 28-JUL-2000; 2000US-221607P.  
XX PR 02-MAY-2001; 2001US-287724P.  
XX  
XX PA (COMP-) COMPUGEN INC.  
XX  
XX PI Shoshan A, Wasserman A, Mintz E, Mintz L, Faigler S;  
XX  
XX DR WPI; 2002-257383/30.  
XX  
XX PT New oligonucleotide libraries comprising oligonucleotides which  
XX selectively hybridize to mRNAs transcribed from a transcription unit of  
XX a genome, useful for detecting tissue-, pathology-, and  
XX developmental-specific genes  
XX  
XX PS Example 1; SEQ ID 25542; 47pp; English.  
XX  
XX CC The present invention describes oligonucleotide libraries for detecting  
XX messenger RNAs that populate a (sub-)transcriptome, where the  
XX (sub-)transcriptome comprises messenger RNAs transcribed from multiple

CC transcription units that populate a genome. The library comprises  
CC several oligonucleotides, each capable of hybridizing selectively to a  
CC set of messenger RNAs transcribed from a given transcription unit of  
CC the genome, which encodes one or more messenger RNA splice variants.  
CC The oligonucleotide libraries are useful for detecting mRNAs from a  
CC biological sample, in expression profiling studies, in qualitatively or  
CC quantitatively characterizing the corresponding transcriptome, and in  
CC detecting RNA transcripts and splice variants of human or animal  
CC transcriptomes. The libraries may also be used as specialised mini  
CC libraries to detect transcripts of a sub-transcriptome under a  
CC particular biological or pathological state, and so allowing the  
CC detection of tissue- and pathology-specific genes such as those genes  
CC only expressed in specific tissue under a specific pathological  
CC condition; to detect developmental specific genes; and to detect RNA  
CC transcripts and splice variants of a transcriptome of a patient suffering  
CC from a particular disorder. ABA27253 to ABA59589 represent  
CC oligonucleotide sequences from rats, humans and mice, which are used in  
CC the exemplification of the present invention.  
CC N.B. The sequence data for this patent did not form part of the printed  
CC specification, but was obtained in electronic format directly from WIPO  
CC at [ftp.wipo.int/pub/published\\_pct\\_sequences](http://ftp.wipo.int/pub/published_pct_sequences).  
XX  
SQ Sequence 65 BP; 8 A; 11 C; 23 G; 23 T; 0 other;  
Query Match 83.3%; Score 15; DB 24; Length 65;  
Best Local Similarity 100.0%; Pred. No. 21;  
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
OY 1 GGCTGGTGTACCTG 15  
Db 8 GGCTGGTGTACCTG 22  
RESULT 20  
ABV56544  
XX ABEV56544 standard; cDNA; 609 BP.  
XX  
AC ABEV56544;  
XX  
DT 17-SEP-2002 (first entry)  
XX  
DE Human prostate expression marker cDNA 56535.  
XX  
XX Human: prostate cancer; cytostatic; carcinogen; pharmacodynamic marker;  
XX pharmacogenomic marker; gene; ss.  
XX  
XX OS Homo sapiens.  
XX  
XX PN WO200160860-A2.  
XX  
XX PD 23-AUG-2001.  
XX  
XX PF 20-FEB-2001; 2001WO-US05171.  
XX  
XX PR 17-FEB-2000; 2000US-183319P.  
XX PR 16-MAR-2000; 2000US-189862P.  
XX PR 25-MAY-2000; 2000US-207454P.  
XX PR 08-JUN-2000; 2000US-211314P.  
XX PR 18-JUL-2000; 2000US-219007P.  
XX PR 13-DEC-2000; 2000US-255281P.  
XX  
XX PA (MILL-) MILLENNIUM PREDICTIVE MEDICINE INC.  
XX  
XX PI Schlegel R, Endege WO, Monahan JB;  
XX  
XX DR WPI; 2001-662795/76.  
XX  
XX PT Novel isolated nucleic acid molecule associated with cancerous state of  
XX prostate cells and correlating with presence of prostate cancer, useful  
XX for detecting presence of prostate cancer. Stage of prostate cancer  
XX  
XX PS Claim 1; Page 10906; 11750pp; English.



CC The invention relates to an isolated nucleic acid molecule (1) comprising  
 CC a nucleotide sequence given in Tables 1-9 (ABV00010-ABV62213) of the  
 CC specification or its complement. (1) is useful for:  
 CC (a) assessing whether a patient is afflicted with prostate cancer;  
 CC (b) monitoring the progression of prostate cancer in a patient;  
 CC (c) assessing the efficacy of a test compound to inhibit prostate  
 CC cancer in a patient;  
 CC (d) assessing the efficacy of a therapy for inhibiting prostate cancer  
 CC in a patient;  
 CC (e) selecting a composition for inhibiting prostate cancer in a patient;  
 CC (f) assessing the prostate cell carcinogenic potential of a compound;  
 CC (g) determining whether prostate cancer has metastasized in a patient;  
 CC (h) assessing the aggressiveness or indolence of prostate cancer in a  
 CC patient;  
 CC (i) is also useful as a pharmacodynamic or pharmacogenomic marker.

SQ Sequence 609 BP; 186 A; 116 C; 99 G; 208 T; 0 other;

Query Match 83.3%; Score 15; DB 23; Length 609;  
 Best Local Similarity 100.0%; Pred. No. 21;  
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 CTGGTGTCACTGTT 17  
 |||||  
 DB 45 CTGGTGTCACTGTT 59

RESULT 21  
 AAF1311  
 ID AAF1311 standard; cDNA; 616 BP.  
 AC AAF1311;  
 XX  
 DT 13-MAR-2001 (first entry)  
 XX  
 DE Aspergillus niger EST SEQ ID NO:3834.  
 XX  
 DE Multiple gene expression; filamentous fungal cell; EST;  
 XX expressed sequence tag; Fusarium venenatum; Aspergillus niger;  
 KM Aspergillus oryzae; Trichoderma reesei; identification; recombination;  
 KM culture condition; environmental stress; spore morphogenesis;  
 KM metabolic pathway engineering; catabolic pathway engineering; ss.  
 XX  
 OS Aspergillus niger.  
 XX  
 PN WO200056762-A2.  
 XX  
 PD 28-SEP-2000.  
 XX  
 PF 22-MAR-2000; 2000MO-US07781.  
 XX  
 PR 22-MAR-1999; 99US-0273623.  
 XX  
 PA (NOVO) NOVO NORDISK BIOTECH INC.  
 PA (NOVO) NOVO NORDISK AS.  
 XX  
 PI Berka RM, Rey MM, Shuster JR, Kaupinen S, Clausen IG, Olsen PB;  
 XX  
 DR WPI; 2000-594572/56.  
 XX  
 PT Monitoring differential expression of genes in filamentous fungal cells  
 PT uses fluorescence-labeled nucleic acids isolated from the cells and a  
 PT substrate of expressed sequence tags -  
 XX  
 PS Claim 87; Page 1716; 3161pp; English.  
 XX  
 CC The present invention describes a method for monitoring differential  
 CC expression of genes in a first filamentous fungal (PF) cell relative to  
 CC expression of the same genes in one or more second filamentous fungal  
 CC cells. The method uses fluorescence-labeled nucleic acids isolated from  
 CC the PF cells and a substrate of expressed sequence tags (EST). The ESTs  
 CC are used in the methods for monitoring differential expression of genes  
 CC in a first filamentous fungal (PF) cell relative to expression of the

CC same genes in one or more second filamentous fungal cells. Monitoring  
 CC the global expression of genes from PF cells allows the production  
 CC potential of the microorganisms to be improved. New genes may be  
 CC discovered, possible functions of unknown open reading frames can be  
 CC identified and gene copy number variation and stability can be  
 CC monitored. The expression of genes can be used to study how PF cells  
 CC adapt to changes in culture conditions, environmental stress, spore  
 CC morphogenesis, recombination, metabolic or catabolic pathway  
 CC engineering. Using ESTs provides several advantages over genomic or  
 CC random cDNA clones including elimination of redundancy as one spot on an  
 CC array equals one gene or open reading frame, and organisation of the  
 CC microarrays based on function of the gene products to facilitate  
 CC analysis of the results. AAF07478 to AAF11247 represents ESTs from  
 CC Fusarium venenatum; AAF11248 to AAF11853 represents ESTs from  
 CC Aspergillus niger; AAF1854 to AAF14878 represents ESTs from Aspergillus  
 CC niger; AAF14879 to AAF15337 represents ESTs from Trichoderma reesei, which are  
 CC all specifically claimed in the present invention.

SQ Sequence 616 BP; 138 A; 179 C; 141 G; 146 T; 12 other;

Query Match 83.3%; Score 15; DB 21; Length 616;  
 Best Local Similarity 100.0%; Pred. No. 21;  
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGCTGGTGTCACTG 15  
 |||||  
 DB 244 GGCTGGTGTCACTG 258

RESULT 22  
 AAH45451  
 ID AAH45451 standard; cDNA; 1419 BP.  
 AC AAH45451;  
 XX  
 DT 06-SEP-2001 (first entry)  
 XX  
 DE Murine epilepsy-causing gene Epm2a cDNA sequence.  
 XX  
 DE Mouse; Epm2a; epilepsy; transgenic animal; knockout mouse; ss.  
 XX  
 OS Mus musculus.  
 XX  
 FH Key Location/Qualifiers  
 FT CDS 67..1059  
 FT /tag= a  
 FT /product= "Epm2a"  
 FT /note= "Protein tyrosine phosphatase"  
 XX  
 PN JP2001095587-A.  
 XX  
 PD 10-APR-2001.  
 XX  
 PF 01-OCT-1999; 99JP-0281632.  
 XX  
 PR 01-OCT-1999; 99JP-0281632.  
 XX  
 PA (RIKA) RIKAGAKU KENKYUSHO.  
 XX  
 DR WPI; 2001-341250/36.  
 DR P-PDB; AAG62454.  
 XX  
 PT Mouse epilepsy-causing gene for the analysis of epilepsy and use in a  
 PT mouse model of epilepsy -  
 XX  
 PS Claim 2; Page 6-8; 10pp; Japanese.  
 XX  
 CC This invention relates to a murine gene (Epm2a) and its encoded protein,  
 CC which cause epilepsy. Epm2a is a protein tyrosine phosphatase. Included  
 CC in the invention is a method of preparing an Epm2a knockout mouse. The  
 CC gene can be used for the analysis of Epm2a expression and to create an  
 CC epilepsy model animal. The present sequence represents cDNA encoding  
 CC murine epilepsy causing Epm2a protein.

XX SQ Sequence 1419 BP; 293 A; 376 C; 437 G; 313 T; 0 other;  
Query Match 83.3%; Score 15; DB 22; Length 1419;  
Best Local Similarity 100.0%; Pred. No. 21;  
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
1 GGCTGTCACCTG 15  
Db 1146 GGCTGTCACCTG 1160  
RESULT 23  
ABZ11825  
ID ABZ11825 standard; cDNA; 1699 BP.  
XX AC ABZ11825;  
XX DT 20-JAN-2003 (first entry)  
XX DE Human polynucleotide SEQ ID NO 707.  
XX KW Human; genome mapping; gene therapy; food supplement; virus; fungus;  
KW cell-proliferative disorder; neurodegenerative disease; bacterial;  
KW Parkinson's disease; Alzheimer's disease; autoimmune disease;  
KW multiple sclerosis; diabetes; genetic disorder; wound; burn; infection;  
KW arthritis; cytotoxic; immunomodulator; nootropic; neuroprotective;  
KW antiparkinsonian; antidiabetic; immunosuppressive; dermatological;  
KW haemostatic; vulnery; fungicide; antibacterial; virucide; protozoacide;  
KW antithratic; gene; ss.  
XX OS Homo sapiens.  
XX PN WO200270539-A2.  
XX PD 12-SEP-2002.  
XX PF 05-MAR-2002; 2002MO-US05095.  
XX PR 05-MAR-2001; 2001US-0799451.  
XX PA (HYSE-) HYSEQ INC.  
XX PI Tang YT, Zhou P, Goodrich RM, Asundi V, Zhang J, Zhao QA, Ren F,  
PI Xue AJ, Yang Y, Ma Y, Yamazaki V, Chen R, Wang Z, Ghosh M,  
PI Wehrman T, Wang J, Wang D, Drmanac RT;  
XX DR WPI; 2002-759812/82.  
XX DR P-PSDB; ABP69608.  
XX PT New polynucleotides comprising sequences assembled from expressed  
PT sequence tags (ESTs), useful for treating cell-proliferative,  
PT neurodegenerative, autoimmune, genetic, myeloid or lymphoid, or  
PT platelet or coagulation disorders -  
XX PS Claim 1; SEQ ID NO 707; 1012pp + Sequence Listing: English.  
XX CC The invention relates to an isolated polynucleotide (I) comprising a  
CC nucleotide sequence selected from any of 948 sequences  
CC (ABZ1119-ABZ12066) or their mature protein coding portion, active domain  
CC coding protein or complementary sequences. The polynucleotides are useful  
CC for identifying expressed genes or for physical mapping of human genome.  
CC The encoded polypeptides (ABP68902-ABP69849) are useful as molecular  
CC weight markers, as a food supplement, for generating antibodies, in  
CC medical imaging, screening and diagnostic assays and for treating  
CC cell-proliferative disorders (cancer), neurodegenerative diseases  
CC (Parkinson's or Alzheimer's disease), autoimmune diseases (multiple  
CC sclerosis, diabetes, lupus) genetic disorders, myeloid or lymphoid  
CC disorders, platelet or coagulation disorders, wound, burns, incision,  
CC ulcers, liver or lung fibrosis, infections (bacterial, viral, fungal,  
CC parasitic), arthritis, etc.  
CC Note: The sequence data for this patent did not form part of the printed  
CC specification, but was obtained in electronic format directly from WIPO

CC at ftp.wipo.int/pub/published\_pct\_sequences.  
XX SQ Sequence 1699 BP; 512 A; 296 C; 287 G; 604 T; 0 other;  
Query Match 83.3%; Score 15; DB 24; Length 1699;  
Best Local Similarity 100.0%; Pred. No. 21;  
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
3 CTGCTGTCACCTGT 17  
Db 742 CTGCTGTCACCTGT 756  
RESULT 24  
AAV02313  
ID AAV02313 standard; cDNA; 1808 BP.  
XX AC AAV02313;  
XX DT 06-MAY-1998 (first entry)  
XX DE C16N gene for promoting neuron survival and type I collagen production.  
XX KW C16; C16N; neuron survival; type I collagen; calcium regulation;  
KW hypercalcaemia; hypertension; diabetes; arteriosclerosis; cancer;  
KW myocardial infarction; hydroxyapatite; osteoblast; ds.  
XX OS Mus sp.  
XX FH Key Location/Qualifiers  
XX FT CDS 1..1740  
XX FT /tag= a  
XX FT /product= "C16N"  
XX PN WO9740150-A1.  
XX PD 30-OCT-1997.  
XX PF 23-APR-1997; 97WO-JP01391.  
XX PR 10-FEB-1997; 97JP-0041562.  
XX PR 23-APR-1996; 96JP-0127954.  
XX PA (SUMU) SUMITOMO PHARM CO LTD.  
XX PI Ishiduka Y, Mochizuki R;  
XX DR WPI; 1997-535834/49.  
XX DR P-PSDB; AAW31366.  
XX PT Proteins C16 and C16N promote neuron survival and type I collagen  
PT production - for treatment of diseases involving collagen  
PT production, calcium regulation or neuron survival  
XX PS Claim 6; Page 61-62; 86pp; Japanese.  
XX CC The present sequence encodes C16N which can: (a) induce differentiation  
CC into cells which can degrade hydroxyapatite; (b) maintain neuron  
CC survival; (c) inhibit osteoblast proliferation; and (d) promote type I  
CC collagen expression in osteoblasts. C16 and C16N are agents for the  
CC treatment of a broad range of diseases including hypercalcaemia,  
CC hypertension, diabetes, arteriosclerosis, myocardial infarction and  
CC terminal cancer. They may also be used as a screen for potential  
CC inhibitors of their activity for possible medicinal use. Transgenic  
CC animals containing DNA coding for the proteins can be used as model  
CC organisms and for the production of recombinant C16/C16N.  
XX SQ Sequence 1808 BP; 336 A; 548 C; 551 G; 373 T; 0 other;  
Query Match 83.3%; Score 15; DB 18; Length 1808;  
Best Local Similarity 100.0%; Pred. No. 21;  
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 GGCTGGTGTACCTG 15  
|||  
XX 949 GGCTGGTGTACCTG 963

RESULT 25  
AAZ44730  
ID AAZ44730 standard; cDNA; 2293 BP.

AC AAZ44730;

DT 17-APR-2000 (first entry)

DE Human C16N-1 cDNA.

KM C16N; C16N-1; C16N-2; human; cartilage cell differentiation; therapy;  
cartilage disorder treatment agent; arthritis; rheumatoid arthritis; ss.

OS Homo sapiens.

PH Key Location/Qualifiers

FT CDS 107..2293

FT /tag= a

FT /product= "C16N-1"

PN WO200001405-A1.

PD 13-JAN-2000.

PF 02-JUL-1999; 99WO-JP03577.

PR 06-JUL-1998; 98JP-0190889.

PA (SUMU ) SUMITOMO PHARM CO LTD.

PI Ishiduka Y, Mochizuki R;

DR WPI: 2000-126903/11.

XX P-PSDB; AAY51326.

XX Cartilage cell differentiation promoters or remedies for cartilage

XX failure

XX Example 2; Page 36-41; 55pp; Japanese.

CC This invention describes a novel cartilage cell differentiation  
CC promoter, or cartilage disorder treatment agent, which contains C16N,  
CC C16N-1, C16N-2, or their protein analogs, or one of the genes encoding  
CC them, as active ingredient. The promoters and cartilage disorder  
CC treatment agents are for the therapy of cartilage failure including  
CC deformans arthritis, diseases due to cartilage formation abnormality,  
CC or damage to cartilage in bone traction, joint cartilage caused by injury,  
CC arthritis, spondylosis arthritis, chronic rheumatoid arthritis, tuberculous  
CC fever, systemic erythematous, deformative spinal diseases, and  
CC intervertebral hernia. This sequence encodes the human C16N-1 protein  
CC which is described in the method of the invention.

XX Sequence 2293 BP; 417 A; 716 C; 691 G; 469 T; 0 other;

XX Query Match 83.3%; Score 15; DB 21; Length 2293;

XX Best Local Similarity 100.0%; Pred. No. 21;

XX Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 GGCTGGTGTACCTG 15

|||  
XX 1505 GGCTGGTGTACCTG 1519

RESULT 26  
AAZ40179  
ID AAZ40179 standard; cDNA; 2293 BP.

XX AAZ40179;  
AC 22-FEB-2000 (first entry)

DT Human C16N-1 coding sequence.

DE C16N-1; C16N-2; human; myeloid cell differentiation; neuron;

KM hydroxyapatite decomposition; osteoblast growth inhibitor;

XX type I collagen expression; ds.

OS Homo sapiens.

PN JP11308995-A.

PD 09-NOV-1999.

PF 28-APR-1998; 98JP-0134440.

PR 28-APR-1998; 98JP-0134440.

PA (SUMU ) SUMITOMO SEIYAKU KK.

PI WPI: 2000-046933/04.

DR P-PSDB; AAY55017.

XX New proteins C16N-1 and C16N-2 or a gene coding them - can have

XX activity limited to specific tissue or site

XX Claim 2; Page 18-21; 30pp; Japanese.

CC This sequence encodes a C16N-1 protein of the invention. The invention  
CC also relates to C16N-2 proteins. The proteins of the invention have the  
CC following features: (1) it has a differentiation inducing activity,  
CC inducing differentiation of a myeloid cell to a cell having  
CC hydroxyapatite decomposing activity; (2) it has an activity of  
CC maintaining the survival of neuron; (3) it has an activity of inhibiting  
CC growth of osteoblasts; and (4) it has an activity of promoting expression  
CC of type I collagen in osteoblasts. The activity of the proteins can be  
CC limited to a specific tissue or a specific site.

XX Sequence 2293 BP; 417 A; 716 C; 691 G; 469 T; 0 other;

XX Query Match 83.3%; Score 15; DB 21; Length 2293;

XX Best Local Similarity 100.0%; Pred. No. 21;

XX Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 GGCTGGTGTACCTG 15

|||  
XX 1505 GGCTGGTGTACCTG 1519

RESULT 27  
AAZ44731  
ID AAZ44731 standard; cDNA; 2301 BP.

AC AAZ44731;

DT 17-APR-2000 (first entry)

DE Human C16N-2 cDNA.

KM C16N; C16N-1; C16N-2; human; cartilage cell differentiation; therapy;

XX cartilage disorder treatment agent; arthritis; rheumatoid arthritis; ss.

OS Homo sapiens.

PH Key Location/Qualifiers

FT CDS 95..2233

FT /tag= a

PN WO200001405-A1.

XX 13-JAN-2000.  
PD 02-JUL-1999; 99WO-JP03577.  
XX 06-JUL-1998; 98UP-0190889.  
XX (SUMU) SUMITOMO PHARM CO LTD.  
XX Ishiduka Y, Mochizuki R;  
XX WPI, 2000-126903/11.  
DR P-PSDB; AAY51327.  
XX  
XX Cartilage cell differentiation promoters or remedies for cartilage  
PT failure \_  
XX  
XX Example 2; Page 45-50; 55pp; Japanese.  
XX  
XX This invention describes a novel cartilage-cell differentiation  
CC promoter, or cartilage disorder treatment agent, which contains C16N,  
CC C16N-1, C16N-2, or their protein analogs, or one of the genes encoding  
CC them, as active ingredient. The promoters and cartilage disorder  
CC treatment agents are for the therapy of cartilage failure including  
CC deformans arthritis, diseases due to cartilage formation abnormality,  
CC deletion of cartilage in bone fraction, joint cartilage caused by injury,  
CC or damage to articular disc, acute purulent arthritis, tuberculous  
CC arthritis, sphyulous arthritis, chronic rheumatoid arthritis, rheumatic  
CC fever, systemic erythematosis, deformative spinal diseases, and  
CC intervertebral hernia. This sequence encodes the human C16N-2 protein  
CC which is described in the method of the invention.  
XX  
SQ Sequence 2301 BP; 425 A; 714 C; 703 G; 459 T; 0 other;  
XX  
XX  
XX Query Match 83.3%; Score 15; DB 21; Length 2301;  
XX Best Local Similarity 100.0%; Pred. No. 21;  
XX Matches 15; Conservativity 0; Mismatches 0; Indels 0; Gaps 0;  
OY 1 GGCTGGTGTACCTG 15  
DB 1442 GGCTGGTGTACCTG 1456  
XX  
XX  
XX RESULT 28  
XX AAZ40180  
XX ID AAZ40180 standard; cDNA; 2301 BP.  
XX  
XX AAZ40180;  
XX  
XX 22-FEB-2000 (first entry)  
XX  
XX Human C16N-2 coding sequence.  
XX  
XX C16N-1; C16N-2; human; myeloid cell differentiation; neuron;  
XX hydroxyapatite decomposition; osteoblast growth inhibitor;  
XX type I collagen expression; ds.  
XX  
XX Homo sapiens.  
XX  
XX JPI1308995-A.  
XX  
XX 09-NOV-1999.  
XX  
XX 28-APR-1998; 98UP-0134440.  
XX  
XX 28-APR-1998; 98UP-0134440.  
XX  
XX (SUMU) SUMITOMO SEIYAKU KK.  
XX  
XX WPI; 2000-046933/04.  
XX P-PSDB; AAY55018.  
XX  
XX New proteins C16N-1 and C16N-2 or a gene coding them - can have

PT activity limited to specific tissue or site  
XX  
XX Claim 2; Page 23-26; 30pp; Japanese.  
XX  
XX This sequence encodes a C16N-2 protein of the invention. The invention  
CC also relates to C16N-1 proteins. The proteins of the invention have the  
CC following features: (1) it has a differentiation inducing activity,  
CC including differentiation of a myeloid cell to a cell having  
CC hydroxyapatite decomposing activity; (2) it has an activity of  
CC maintaining the survival of neuron; (3) it has an activity of inhibiting  
CC growth of osteoblasts; and (4) it has an activity of promoting expression  
CC of type I collagen in osteoblasts. The activity of the proteins can be  
CC limited to a specific tissue or a specific site.  
XX  
SQ Sequence 2301 BP; 425 A; 714 C; 703 G; 459 T; 0 other;  
XX  
XX  
XX Query Match 83.3%; Score 15; DB 21; Length 2301;  
XX Best Local Similarity 100.0%; Pred. No. 21;  
XX Matches 15; Conservativity 0; Mismatches 0; Indels 0; Gaps 0;  
OY 1 GGCTGGTGTACCTG 15  
DB 1442 GGCTGGTGTACCTG 1456  
XX  
XX  
XX RESULT 29  
XX AAZ33584  
XX ID AAZ33584 standard; cDNA; 2952 BP.  
XX  
XX AAZ33584;  
XX  
XX 08-DEC-1999 (first entry)  
XX  
XX Human breast tumour-associated EST 44.  
XX  
XX Expressed sequence tag; EST; human; breast; cancer; cytostatic;  
XX KW medicaments; gene therapy; treatment; fat metabolism; ss.  
XX  
XX Homo sapiens.  
XX  
XX DE19813835-A1.  
XX  
XX 23-SEP-1999.  
XX  
XX 20-MAR-1998; 98DE-1013835.  
XX  
XX 20-MAR-1998; 98DE-1013835.  
XX  
XX (META-) METAGEN GES GENOMFORSCHUNG MBH.  
XX  
XX Specht T, Hinzmann B, Schmitt A, Pilarsky C, Dahl E, Rosenthal A;  
XX WPI; 1999-528979/45.  
XX  
XX Human nucleic acid sequences and protein products from normal breast  
XX PT tissue, useful for breast cancer therapy  
XX  
XX Claim 3; 132-133; 206pp; German.  
XX  
XX This invention describes novel human nucleic acid sequences from normal  
CC breast tissue which have cytostatic activity. The nucleic acid sequences  
CC can be used to produce and isolate full-length gene sequences. They can  
CC be used to express proteins, which can be used as tools to find an  
CC activity against breast cancer. The sequences can be used in sense or  
CC antisense form. They are especially useful for medicaments for gene  
CC therapy to treat breast cancer and for treating illnesses associated  
CC with fat metabolism. AAZ33541-233610 represent expressed sequence tags  
CC described in the method of the invention.  
XX  
SQ Sequence 2952 BP; 925 A; 564 C; 582 G; 881 T; 0 other;  
XX  
XX  
XX Query Match 83.3%; Score 15; DB 20; Length 2952;  
XX Best Local Similarity 100.0%; Pred. No. 20;

Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
Query 3 CTGGTGTCACTGTT 17  
|||||  
Db 2836 CTGGTGTCACTGTT 2850

RESULT 30  
AAC77831  
ID AAC77831 standard; cDNA; 3035 BP.

AAC77831;

08-FEB-2001 (first entry)

Human cancer associated gene sequence SEQ ID NO:225.

Human; cancer associated gene; cancer antigen; detection; cancer;  
diagnosis; cytostatic; proliferative; vulnery; immunomodulator;  
antidiabetic; antiaesthetic; antirheumatic; antithrombotic; antiviral;  
antiinflammatory; antihypertensive; antiallergic; antibacterial; cardiant;  
dermatological; neuroprotective; thrombolytic; coagulant; nocotropic;  
vasotropic; antipsoriatic; angiogenic; gene therapy; inflammation;  
immune disorder; haematopoietic cell disorder; autoimmune disorder;  
allergic reaction; graft versus host disease; organ rejection;  
haemostatic; thrombolytic; cardiovascular disorder; infection;  
neurological disease; drug screening; ss.

Homo sapiens.

WO200055350-A1.

21-SEP-2000.

08-MAR-2000; 2000WO-US05882.

12-MAR-1999; 99US-0324270.

(HUMA-) HUMAN GENOME SCI INC.

Rosen CA, Ruben SM;

WPI; 2000-587533/55.

P-PSDB; AAB43622.

Novel isolated nucleic acids comprising sequences encoding peptides  
useful for treating or diagnosing e.g. cancer -

Claim 1; Page 796-797; 2352pp; English.

AAC77607 to AAC78448 encode the human cancer associated proteins given  
in AAB4398 to AAB4429. The proteins can have activities based on the  
tissues and cells the genes are expressed in. Example of activities  
include: cytostatic; proliferative; vulnery; immunomodulator;  
antiinflammatory; antihypertensive; antiallergic; antibacterial;  
antiviral;  
dermatological; neuroprotective; cardiant; thrombolytic; coagulant;  
nocotropic; vasotropic; antipsoriatic and angiogenic. The  
polynucleotides and polypeptides can be used for preventing, treating or  
ameliorating medical conditions and diagnosing pathological conditions.  
Polynucleotides, polypeptides, antibodies, agonists and antagonists from  
the present invention may be used to treat immune disorders by activating  
or inhibiting the proliferation, differentiation or mobilization of  
immune cells, to treat disorders of haematopoietic cells, autoimmune  
disorders, allergic reactions, graft versus host disease and organ  
rejection, modulate haemostatic or thrombolytic activity, modulate  
inflammation, cancers, cardiovascular disorders, neurological disease and  
bacterial or viral infections. The peptides, nucleotides, antibodies,  
agonists and antagonists may be also be used in drug screens. AAC78449 to  
AAC78457 and AAB44240 represent sequences used in the exemplification of  
the present invention.

Sequence 3035 BP; 955 A; 575 C; 588 G; 912 T; 5 other;

Query Match 83.3%; Score 15; DB 21; Length 3035;  
Best Local Similarity 100.0%; Pred. No. 20;  
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Query 3 CTGGTGTCACTGTT 17  
|||||  
Db 2771 CTGGTGTCACTGTT 2785

RESULT 31  
AAH34809  
ID AAH34809 standard; cDNA; 3035 BP.

AAH34809;

03-SEP-2001 (first entry)

Human colon cancer antigen encoding cDNA SEQ ID NO:1891.

Human; colon cancer; colon cancer antigen; diagnosis; detection;  
colorectal carcinoma; chromosome 2; ss.

Homo sapiens.

WO200122920-A2.

05-APR-2001.

28-SEP-2000; 2000WO-US26524.

29-SEP-1999; 99US-0157137.

03-NOV-1999; 99US-0163280.

(HUMA-) HUMAN GENOME SCI INC.

Ruben SM, Barash SC, Birse CE, Rosen CA;

WPI; 2001-235357/24.

P-PSDB; AAG75404.

Nucleic acids encoding 4277 human colon cancer-associated polypeptides,  
useful for preventing, diagnosing and/or treating colorectal cancers -

Claim 1; Page 3405-3406; 9803pp; English.

AAH32943 to AAH37195 and AAG73514 to AAG77788 represent human colon  
cancer-associated nucleic acid molecules (N) and proteins (P), where  
the proteins are collectively known as colon cancer antigens. The colon  
cancer antigens have cytostatic activity and can be used in gene  
therapy and vaccine production. N and P may be used in the prevention,  
diagnosis and treatment of diseases associated with inappropriate P  
expression. For example, N and P may be used to treat disorders  
associated with decreased expression by rectifying mutations or deletions  
in a patient's genome that affect the activity of P by expressing  
inactive proteins or to supplement the patient's own production of P.  
Additionally, N may be used to produce the colon cancer-associated Ps,  
by inserting the nucleic acids into a host cell and culturing the cell  
to express the proteins. N and P can be used in the prevention, diagnosis  
and treatment of colorectal carcinomas and cancers. AAH37196 to AAH37204  
and AAG77789 represent sequences used in the exemplification of the  
present invention.

N.B. Pages 666 to 682 and page 7053 of the sequence listing were  
missing at time of publication, meaning no sequences are present for  
SEQ ID NO:1027 to 1052, 7921 and 7922.

Sequence 3035 BP; 955 A; 575 C; 590 G; 912 T; 3 other;

Query Match 83.3%; Score 15; DB 22; Length 3035;  
Best Local Similarity 100.0%; Pred. No. 20;  
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Query 3 CTGGTGTCACTGTT 17

Db 2771 CTGCTGTACCTGTT 2785

# RESULT 32

AAV02312 standard; cDNA; 3065 BP.

AAV02312;

06-MAY-1998 (first entry)

C16N gene for promoting neuron survival and type 1 collagen production.

C16; C16N; neuron survival; type 1 collagen; calcium regulation;

hypercalcaemia; hypertension; diabetes; arteriosclerosis; cancer;

myocardial infarction; hydroxyapatite; osteoblast; ds.

Mus sp.

Key Location/Qualifiers

CDS 236..1975

FT /tag= a

FT /product= "C16N"

FT

PA MO9740150-A1.

PN 30-OCT-1997.

XX 23-APR-1997; 97WO-JP01391.

XX 10-FEB-1997; 97JP-0041562.

XX 23-APR-1996; 96JP-0127954.

XX (SUMU) SUMITOMO PHARM CO LTD.

PI Ishiduka Y, Mochizuki R;

XX WPI; 1997-535834/49.

XX P-PSDB; AAW31364.

XX Proteins C16 and C16N promote neuron survival and type 1 collagen

XX production - for treatment of diseases involving collagen

XX production, calcium regulation or neuron survival

XX Claim 6; Page 55-57; 86pp; Japanese.

XX The present sequence encodes C16N which can: (a) induce differentiation

XX into cells which can degrade hydroxyapatite; (b) maintain neuron

XX survival; (c) inhibit osteoblast proliferation; and (d) promote type 1

XX collagen expression in osteoblasts. C16 and C16N are agents for the

XX treatment of a broad range of diseases including hypercalcaemia,

XX hypertension, diabetes, arteriosclerosis, myocardial infarction and

XX terminal cancer. They may also be used as a screen for potential

XX inhibitors of their activity for possible medicinal use. Transgenic

XX animals containing DNA coding for the proteins can be used as model

XX organisms and for the production of recombinant C16/C16N.

AAZ44729;

17-APR-2000 (first entry)

Murine C16N-2 cDNA.

C16N; C16N-1; C16N-2; murine; cartilage cell differentiation; therapy;

cartilage disorder treatment agent; arthritis; rheumatoid arthritis; ss.

Mus musculus.

Key Location/Qualifiers

CDS 109..2247

FT /tag= a

FT /product= "C16N-2"

FT

PA MO200001405-A1.

PN 13-JAN-2000.

XX 02-JUL-1999; 99WO-JP03577.

XX 06-JUL-1998; 98JP-0190889.

XX (SUMU) SUMITOMO PHARM CO LTD.

PI Ishiduka Y, Mochizuki R;

XX WPI; 2000-126903/11.

XX P-PSDB; AAY51325.

XX Cartilage cell differentiation promoters or remedies for cartilage

XX failure

XX Example 1; Page 27-33; 55pp; Japanese.

XX This invention describes a novel cartilage cell differentiation

XX promoter, or cartilage disorder treatment agent, which contains C16N,

XX C16N-1, C16N-2, or their protein analogs, or one of the genes encoding

XX them, as active ingredient. The promoters and cartilage disorder

XX treatment agents are for the therapy of cartilage failure including

XX deformans arthritis, diseases due to cartilage formation abnormality,

XX deletion of cartilage in bone fraction, joint cartilage caused by injury,

XX or damage to articular disc, acute purulent arthritis, tuberculous

XX arthritis, synovial arthritis, chronic rheumatoid arthritis, rheumatic

XX fever, systemic erythematous, deformative spinal diseases, and

XX intervertebral hernia. This sequence encodes the murine C16N-2 protein

XX which is described in the method of the invention.

XX Sequence 3337 BP; 633 A; 1037 C; 976 G; 691 T; 0 other;

XX Query Match 83.3%; Score 15; DB 21; Length 3337;

XX Best Local Similarity 100.0%; Pred. No. 20;

XX Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

AAZ40178

22-FEB-2000 (first entry)

Mouse C16N-2 coding sequence.

C16N-1; C16N-2; mouse; myeloid cell differentiation; neuron;

hydroxyapatite decomposition; osteoblast growth inhibitor;

type I collagen expression; ds.

Key Location/Qualifiers

CDS 109..2247

FT /tag= a

FT /product= "C16N-2"

FT

PA MO200001405-A1.

PN 13-JAN-2000.

XX 02-JUL-1999; 99WO-JP03577.

XX 06-JUL-1998; 98JP-0190889.

XX (SUMU) SUMITOMO PHARM CO LTD.

PI Ishiduka Y, Mochizuki R;

XX WPI; 2000-126903/11.

XX P-PSDB; AAY51325.

XX Cartilage cell differentiation promoters or remedies for cartilage

XX failure

XX Example 1; Page 27-33; 55pp; Japanese.

XX This invention describes a novel cartilage cell differentiation

XX promoter, or cartilage disorder treatment agent, which contains C16N,

XX C16N-1, C16N-2, or their protein analogs, or one of the genes encoding

XX them, as active ingredient. The promoters and cartilage disorder

XX treatment agents are for the therapy of cartilage failure including

XX deformans arthritis, diseases due to cartilage formation abnormality,

XX deletion of cartilage in bone fraction, joint cartilage caused by injury,

XX or damage to articular disc, acute purulent arthritis, tuberculous

XX arthritis, synovial arthritis, chronic rheumatoid arthritis, rheumatic

XX fever, systemic erythematous, deformative spinal diseases, and

XX intervertebral hernia. This sequence encodes the murine C16N-2 protein

XX which is described in the method of the invention.

XX Sequence 3337 BP; 633 A; 1037 C; 976 G; 691 T; 0 other;

XX Query Match 83.3%; Score 15; DB 21; Length 3337;

XX Best Local Similarity 100.0%; Pred. No. 20;

XX Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

AAZ40178

22-FEB-2000 (first entry)

Mouse C16N-2 coding sequence.

C16N-1; C16N-2; mouse; myeloid cell differentiation; neuron;

hydroxyapatite decomposition; osteoblast growth inhibitor;

type I collagen expression; ds.

Key Location/Qualifiers

CDS 109..2247

FT /tag= a

FT /product= "C16N-2"

FT

PA MO200001405-A1.

PN 13-JAN-2000.

XX 02-JUL-1999; 99WO-JP03577.

XX 06-JUL-1998; 98JP-0190889.

XX (SUMU) SUMITOMO PHARM CO LTD.

PI Ishiduka Y, Mochizuki R;

XX WPI; 2000-126903/11.

XX P-PSDB; AAY51325.

XX Cartilage cell differentiation promoters or remedies for cartilage

XX failure

XX Example 1; Page 27-33; 55pp; Japanese.

XX This invention describes a novel cartilage cell differentiation

XX promoter, or cartilage disorder treatment agent, which contains C16N,

XX C16N-1, C16N-2, or their protein analogs, or one of the genes encoding

XX them, as active ingredient. The promoters and cartilage disorder

XX treatment agents are for the therapy of cartilage failure including

XX deformans arthritis, diseases due to cartilage formation abnormality,

XX deletion of cartilage in bone fraction, joint cartilage caused by injury,

XX or damage to articular disc, acute purulent arthritis, tuberculous

XX arthritis, synovial arthritis, chronic rheumatoid arthritis, rheumatic

XX fever, systemic erythematous, deformative spinal diseases, and

XX intervertebral hernia. This sequence encodes the murine C16N-2 protein

XX which is described in the method of the invention.

XX Sequence 3337 BP; 633 A; 1037 C; 976 G; 691 T; 0 other;

XX Query Match 83.3%; Score 15; DB 21; Length 3337;

XX Best Local Similarity 100.0%; Pred. No. 20;

XX Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

AAZ40178

22-FEB-2000 (first entry)

Mouse C16N-2 coding sequence.

C16N-1; C16N-2; mouse; myeloid cell differentiation; neuron;

hydroxyapatite decomposition; osteoblast growth inhibitor;

type I collagen expression; ds.

Key Location/Qualifiers

CDS 109..2247

FT /tag= a

FT /product= "C16N-2"

FT

PA MO200001405-A1.

PN 13-JAN-2000.

XX 02-JUL-1999; 99WO-JP03577.

XX 06-JUL-1998; 98JP-0190889.

XX (SUMU) SUMITOMO PHARM CO LTD.

PI Ishiduka Y, Mochizuki R;

XX WPI; 2000-126903/11.

XX P-PSDB; AAY51325.

XX Cartilage cell differentiation promoters or remedies for cartilage

XX failure

XX Example 1; Page 27-33; 55pp; Japanese.

XX This invention describes a novel cartilage cell differentiation

XX promoter, or cartilage disorder treatment agent, which contains C16N,

XX C16N-1, C16N-2, or their protein analogs, or one of the genes encoding

XX them, as active ingredient. The promoters and cartilage disorder

XX treatment agents are for the therapy of cartilage failure including

XX deformans arthritis, diseases due to cartilage formation abnormality,

XX deletion of cartilage in bone fraction, joint cartilage caused by injury,

XX or damage to articular disc, acute purulent arthritis, tuberculous

XX arthritis, synovial arthritis, chronic rheumatoid arthritis, rheumatic

XX fever, systemic erythematous, deformative spinal diseases, and

XX intervertebral hernia. This sequence encodes the murine C16N-2 protein

XX which is described in the method of the invention.

XX Sequence 3337 BP; 633 A; 1037 C; 976 G; 691 T; 0 other;

XX Query Match 83.3%; Score 15; DB 21; Length 3337;

XX Best Local Similarity 100.0%; Pred. No. 20;

XX Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

AAZ40178

22-FEB-2000 (first entry)

Mouse C16N-2 coding sequence.

C16N-1; C16N-2; mouse; myeloid cell differentiation; neuron;

hydroxyapatite decomposition; osteoblast growth inhibitor;

type I collagen expression; ds.

Key Location/Qualifiers

CDS 109..2247

FT /tag= a

FT /product= "C16N-2"

FT

PA MO200001405-A1.

PN 13-JAN-2000.

XX 02-JUL-1999; 99WO-JP03577.

XX 06-JUL-1998; 98JP-0190889.

XX (SUMU) SUMITOMO PHARM CO LTD.

PI Ishiduka Y, Mochizuki R;

XX WPI; 2000-126903/11.

XX P-PSDB; AAY51325.

XX Cartilage cell differentiation promoters or remedies for cartilage

XX failure

XX Example 1; Page 27-33; 55pp; Japanese.

XX This invention describes a novel cartilage cell differentiation

XX promoter, or cartilage disorder treatment agent, which contains C16N,

XX C16N-1, C16N-2, or their protein analogs, or one of the genes encoding

XX them, as active ingredient. The promoters and cartilage disorder

XX treatment agents are for the therapy of cartilage failure including

XX deformans arthritis, diseases due to cartilage formation abnormality,

XX deletion of cartilage in bone fraction, joint cartilage caused by injury,

XX or damage to articular disc, acute purulent arthritis, tuberculous

XX arthritis, synovial arthritis, chronic rheumatoid arthritis, rheumatic

XX fever, systemic erythematous, deformative spinal diseases, and

XX intervertebral hernia. This sequence encodes the murine C16N-2 protein

XX which is described in the method of the invention.

XX Sequence 3337 BP; 633 A; 1037 C; 976 G; 691 T; 0 other;

XX Query Match 83.3%; Score 15; DB 21; Length 3337;

XX Best Local Similarity 100.0%; Pred. No. 20;

XX Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

AAZ40178

22-FEB-2000 (first entry)

Mouse C16N-2 coding sequence.

C16N-1; C16N-2; mouse; myeloid cell differentiation; neuron;

hydroxyapatite decomposition; osteoblast growth inhibitor;

type I collagen expression; ds.

Key Location/Qualifiers

CDS 109..2247

FT /tag= a

FT /product= "C16N-2"

FT

PA MO200001405-A1.

PN 13-JAN-2000.

XX 02-JUL

XX Mus sp.  
OS JP1308995-A.  
XX  
XX  
XX 09-NOV-1999.  
XX  
XX 28-APR-1998; 98BP-0134440.  
XX  
XX 28-APR-1998; 98BP-0134440.  
XX  
XX (SUMU ) SUMITOMO SEIYAKU KK.  
XX  
XX WPI; 2000-046933/04.  
XX P-PSDB; AAY55016.  
XX  
XX New proteins Cl6N-1 and Cl6N-2 or a gene coding them - can have  
PT activity limited to specific tissue or site  
XX  
XX Claim 2; Page 13-16; 30pp; Japanese.  
XX  
XX This sequence encodes a Cl6N-2 protein of the invention. The invention  
CC also relates to Cl6N-1 proteins. The proteins of the invention have the  
CC following features: (1) it has a differentiation inducing activity,  
CC inducing differentiation of a myeloid cell to a cell having  
CC hydroxyapatite decomposing activity; (2) it has an activity of  
CC maintaining the survival of neuron; (3) it has an activity of inhibiting  
CC growth of osteoblasts; and (4) it has an activity of promoting expression  
CC of type I collagen in osteoblasts. The activity of the proteins can be  
CC limited to a specific tissue or a specific site.  
XX  
XX Sequence 3337 BP; 633 A; 1037 C; 976 G; 691 T; 0 other;  
SQ

Query Match 83.3%; Score 15; DB 21; Length 3337;  
Best Local Similarity 100.0%; Pred. No. 20;  
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
OY 1 GGCTGGTGCACCTG 15  
|||  
DB 1456 GGCTGGTGCACCTG 1470

RESULT 35  
AAZ44728  
ID AAZ44728 standard; cDNA; 3674 BP.  
XX  
XX AAZ44728;  
XX  
XX 17-APR-2000 (first entry)  
XX  
XX Murine Cl6N-1 cDNA.  
XX  
XX  
XX Cl6N; Cl6N-1; Cl6N-2; murine; cartilage cell differentiation; therapy;  
KW cartilage disorder treatment agent; arthritis; rheumatoid arthritis; ss.  
XX  
XX Mus musculus.  
OS  
XX  
XX Key Location/Qualifiers  
FT CDS 395..2584  
FT /\*tag= a  
FT /product= "Cl6N-1"  
XX  
XX WO200001405-A1.  
XX  
XX 13-JAN-2000.  
XX  
XX 02-JUL-1999; 99WO-JP03577.  
XX  
XX 06-JUL-1998; 98JP-0190889.  
XX  
XX (SUMU ) SUMITOMO PHARM CO LTD.  
XX  
XX Ishiduka Y, Mochizuki R;  
XX  
XX

XX WPI; 2000-126903/11.  
DR P-PSDB; AAY51324.  
XX  
XX Cartilage cell differentiation promoters or remedies for cartilage  
PT failure  
XX  
XX Example 2; Page 18-24; 55pp; Japanese.  
XX

CC This invention describes a novel cartilage cell differentiation  
CC promoter, or cartilage disorder treatment agent, which contains Cl6N,  
CC Cl6N-1, Cl6N-2, or their protein analogs, or one of the genes encoding  
CC them, as active ingredient. The promoters and cartilage disorder  
CC treatment agents are for the therapy of cartilage failure including  
CC deformant arthritis, diseases due to cartilage formation abnormality,  
CC deletion of cartilage in bone fraction, joint cartilage caused by injury,  
CC or damage to articular disc, acute purulent arthritis, tuberculous  
CC arthritis, syphilous arthritis, chronic rheumatoid arthritis, rheumatic  
CC fever, systemic erythematosis, deformative spinal diseases, and  
CC intervertebral hernia. This sequence encodes the murine Cl6N-1 protein  
CC which is described in the method of the invention.

XX Sequence 3674 BP; 695 A; 1146 C; 1044 G; 789 T; 0 other;  
SQ  
Query Match 83.3%; Score 15; DB 21; Length 3674;  
Best Local Similarity 100.0%; Pred. No. 20;  
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
OY 1 GGCTGGTGCACCTG 15  
|||  
DB 1793 GGCTGGTGCACCTG 1807

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Job time : 116.4 secs

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GenCore version 5.1.6  
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: August 14, 2003, 21:44:31 ; Search time 32.25 Seconds

(without alignments)  
273.726 Million cell updates/sec

Title: US-10-074-620-2

Perfect score: 20

Sequence: 1 ccttagagagcaacgccc 20

Scoring table: OLIGO\_NUC

Searched: 559978 seqs, 220691566 residues

Word size: 0

Total number of hits satisfying chosen parameters: 1139956

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Listing first 120 summaries

Database:

Issued Patents\_NA: \*  
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Pred. No. is the number of results predicted by chance to have a  
score greater than or equal to the score of the result being printed,  
and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	20	100.0	120	1	US-08-197-791-29
2	20	100.0	3833	1	US-08-917-130-18
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4	20	100.0	5931	3	US-08-783-774-1
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6	16	80.0	6572	4	US-09-620-312D-823
7	14	70.0	854	4	US-09-439-313-354
8	14	70.0	854	4	US-09-352-616A-354
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18	14	70.0	4249	1	US-08-488-011B-21
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24	13	65.0	126	4	US-08-857-046A-9
25	13	65.0	126	4	US-09-573-252-9
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27	13	65.0	243	3	US-08-756-849-23

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C 31	13	65.0	1525	5	PCT-US92-10284-4	Sequence 4, Appl1
C 32	13	65.0	1645	5	PCT-US94-01321-9	Sequence 9, Appl1
C 33	13	65.0	1970	4	US-09-389-956-79	Sequence 79, Appl1
C 34	13	65.0	2517	4	US-10-020-079-39	Sequence 39, Appl1
C 35	13	65.0	2556	4	US-10-020-079-37	Sequence 37, Appl1
C 36	13	65.0	2592	4	US-10-020-079-31	Sequence 31, Appl1
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C 38	13	65.0	2836	4	US-10-020-079-35	Sequence 35, Appl1
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C 47	13	65.0	176373	3	US-09-128-155-17	Sequence 17, Appl1
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C 57	12	60.0	284	3	US-08-906-791-7	Sequence 7, Appl1
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C 61	12	60.0	323	2	US-08-506-864A-4	Sequence 4, Appl1
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C 63	12	60.0	354	4	US-09-702-705-1618	Sequence 1618, Ap
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C 66	12	60.0	421	3	US-09-265-628-12	Sequence 12, Appl1
C 67	12	60.0	421	3	US-09-001-141-10	Sequence 10, Appl1
C 68	12	60.0	421	3	US-09-653-403-13	Sequence 13, Appl1
C 69	12	60.0	421	4	US-09-643-597-326	Sequence 326, Appl1
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C 81	12	60.0	627	3	US-09-385-982-201	Sequence 201, Appl1
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C 83	12	60.0	669	4	US-09-465-901-19	Sequence 19, Appl1
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C 96	12	60.0	914	4	US-09-177-650-123	Sequence 123, Appl1
C 97	12	60.0	941	4	US-09-312-283C-351	Sequence 351, Appl1
C 98	12	60.0	999	1	US-08-469-649-1	Sequence 1, Appl1
C 99	12	60.0	999	4	US-09-347-878-59	Sequence 59, Appl1
C 100	12	60.0	1016	2	US-08-930-617-1	Sequence 1, Appl1

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102 12 60.0 1049 4 US-09-280-116-176 Sequence 176, App  
103 12 60.0 1056 1 US-08-402-217A-1 Sequence 1, Appl  
104 12 60.0 1056 1 US-08-700-178-1 Sequence 1, Appl  
105 12 60.0 1056 3 US-08-995-654-1 Sequence 1282, Ap  
106 12 60.0 1059 4 US-09-107-532A-1282 Sequence 2348, Ap  
107 12 60.0 1164 4 US-09-107-532A-2348 Sequence 555, App  
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111 12 60.0 1190 4 US-09-390-207-1 Sequence 1, Appl  
112 12 60.0 1207 3 US-09-264-419C-1 Sequence 1980, Ap  
113 12 60.0 1248 4 US-09-134-001C-1980 Sequence 8, Appl  
114 12 60.0 1260 4 US-09-206-166-8 Sequence 50, Appl  
115 12 60.0 1325 2 US-08-632-470-50 Sequence 2, Appl  
116 12 60.0 1339 1 US-07-936-163-2 Sequence 3, Appl  
117 12 60.0 1353 1 US-08-756-299-3 Sequence 3, Appl  
118 12 60.0 1353 2 US-08-964-494-3 Sequence 1, Appl  
119 12 60.0 1380 1 US-07-936-163-1 Patent No. 5463025  
120 12 60.0 1392 6 5463025-2

## ALIGNMENTS

RESULT 1  
US-08-197-791-29  
Sequence 29, Application US/08197791  
Patent No. 5463025-2  
GENERAL INFORMATION:  
APPLICANT: Sorige, Joseph A.  
APPLICANT: Mullinax, Rebecca L.  
TITLE OF INVENTION: NOVEL POLYMERASE COMPOSITIONS AND USES  
TITLE OF INVENTION: THEREOF  
NUMBER OF SEQUENCES: 44  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Limbach and Limbach  
STREET: 2001 Ferry Building  
CITY: San Francisco  
STATE: CA  
COUNTRY: USA  
ZIP: 94111  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: Patentin Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/197,791  
FILING DATE:  
CLASSIFICATION: 424  
PRIOR APPLICATION NUMBER:  
APPLICATION NUMBER: US 08/164,290  
FILING DATE: 08-DEC-1993  
ATTORNEY/AGENT INFORMATION:  
NAME: Bortner, Scott R.  
REGISTRATION NUMBER: 34,298  
REFERENCE/DOCKET NUMBER: STRG 20270 USA  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 415-433-4150  
TELEFAX: 415-433-8716  
INFORMATION FOR SEQ ID NO: 29:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 20 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
MOLECULE TYPE: DNA (genomic)  
HYPOTHETICAL: NO  
ANTI-SENSE: NO  
US-08-197-791-29  
Query Match 100.0%; Score 20; DB 1; Length 20;

Best Local Similarity 100.0%; Pred. No. 0.012;  
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 1 CCTTAGGAGGAACAAGTCCC 20  
Db 1 CCTTAGGAGGAACAAGTCCC 20

RESULT 2  
US-08-917-320-18  
Sequence 18, Application US/08917320  
Patent No. 5824508  
GENERAL INFORMATION:  
APPLICANT: Spaete, Richard and Jackman, Winthrop, T.  
TITLE OF INVENTION: No. 5824508 Splicing Variants of gp350/220  
NUMBER OF SEQUENCES: 18  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Cooley Godward Castro Huddleson & Tatum  
STREET: 5 Palo Alto Square  
CITY: Palo Alto  
STATE: California  
COUNTRY: USA  
ZIP: 94306  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: Patent In Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/917,320  
FILING DATE: 25-AUG-1997  
CLASSIFICATION: 435  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 08/229,291  
FILING DATE: April 18, 1994  
ATTORNEY/AGENT INFORMATION:  
NAME: Luann Geert  
REGISTRATION NUMBER: 31,822  
REFERENCE/DOCKET NUMBER: AVIR-003/00US  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 415-843-5163  
TELEFAX: 415-857-0663  
TELEX: 380816 COOLEYPA  
INFORMATION FOR SEQ ID NO: 18:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 3833 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: double  
TOPOLOGY: unknown  
MOLECULE TYPE: CDNA  
FEATURE:  
NAME/KEY: CDS  
LOCATION: 1014..3734  
US-08-917-320-18  
Query Match 100.0%; Score 20; DB 1; Length 3833;  
Best Local Similarity 100.0%; Pred. No. 0.0089;  
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 1 CCTTAGGAGGAACAAGTCCC 20  
Db 2899 CCTTAGGAGGAACAAGTCCC 2918  
RESULT 3  
PCT-US95-04611A-18  
Sequence 18, Application PC/TUS9504611A  
GENERAL INFORMATION:  
APPLICANT: Spaete, Richard and Jackman, Winthrop, T.  
TITLE OF INVENTION: Non Splicing Variants of gp350/220  
NUMBER OF SEQUENCES: 19  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Cooley Godward Castro Huddleson & Tatum

STREET: 5 Palo Alto Square  
CITY: Palo Alto  
STATE: California  
COUNTRY: USA  
ZIP: 94306  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: Patent In Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: PCT/US95/04611A  
FILING DATE:  
CLASSIFICATION:  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 08/229,291  
FILING DATE: April 18, 1994  
CLASSIFICATION:  
ATTORNEY/AGENT INFORMATION:  
NAME: Luann Geert  
REGISTRATION NUMBER: 31,822  
REFERENCE/DOCKET NUMBER: AVIR-003/00US  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 415-843-5163  
TELEFAX: 415-857-0663  
TELEX: 380816 COOLEYPA  
INFORMATION FOR SEQ ID NO: 18:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 3833 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: double  
TOPOLOGY: unknown  
MOLECULE TYPE: CDNA  
FEATURE:  
NAME/KEY: CDS  
LOCATION: 1014..3734  
PCT-US95-04611A-18  
Query Match 100.0%; Score 20; DB 5; Length 3833;  
Best Local Similarity 100.0%; Pred. NO. 0.0089;  
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
Qy 1 CCTTAGGAGGAACAAGTCCC 20  
Db 2899 CCTTAGGAGGAACAAGTCCC 2918  
RESULT 4  
US-08-783-774-1  
Sequence 1, Application US/08783774  
Patent No. 6054130  
GENERAL INFORMATION:  
APPLICANT: Spaete, Richard  
APPLICANT: Jackman, Winthrop  
TITLE OF INVENTION: NON-SPLICING VARIANTS OF  
TITLE OF INVENTION: GP350/220  
NUMBER OF SEQUENCES: 19  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Pennie & Edmonds  
STREET: 1155 Avenue of the Americas  
CITY: New York  
STATE: NY  
COUNTRY: USA  
ZIP: 10036/2711  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Diskette  
COMPUTER: IBM Compatible  
OPERATING SYSTEM: DOS  
SOFTWARE: FastSeq Version 2.0  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/783,774  
FILING DATE: 15-JAN-1997  
CLASSIFICATION: 435

ATTORNEY/AGENT INFORMATION:  
NAME: Coruzzi, Laura A.  
REGISTRATION NUMBER: 30,742  
REFERENCE/DOCKET NUMBER: 7682-037  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 212-790-9090  
TELEFAX: 212-869-8864  
TELEX: 66141 PENNIE  
INFORMATION FOR SEQ ID NO: 1:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 5931 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
MOLECULE TYPE: DNA  
FEATURE:  
NAME/KEY: Coding Sequence  
LOCATION: 1014...3734  
OTHER INFORMATION:  
US-08-783-774-1  
Query Match 100.0%; Score 20; DB 3; Length 5931;  
Best Local Similarity 100.0%; Pred. NO. 0.0087;  
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
Qy 1 CCTTAGGAGGAACAAGTCCC 20  
Db 2899 CCTTAGGAGGAACAAGTCCC 2918  
RESULT 5  
US-09-556-706B-1  
Sequence 1, Application US/09556706B  
Patent No. 6458364  
GENERAL INFORMATION:  
APPLICANT: Spaete, Richard  
APPLICANT: Jackman, Winthrop  
TITLE OF INVENTION: NON SPLICING VARIANTS OF GP350/220  
FILE REFERENCE: 7682-050-999  
CURRENT APPLICATION NUMBER: US/09/556,706B  
PRIOR FILING DATE: 2000-04-24  
PRIOR FILING DATE: 1997-01-15  
PRIOR APPLICATION NUMBER: 08/783,774  
PRIOR FILING DATE: 1994-04-18  
NUMBER OF SEQ ID NOS: 19  
SOFTWARE: PatentIn version 3.0  
SEQ ID NO 1  
LENGTH: 5931  
TYPE: DNA  
ORGANISM: Virus  
FEATURE:  
OTHER INFORMATION: GP350/220  
US-09-556-706B-1  
Query Match 100.0%; Score 20; DB 4; Length 5931;  
Best Local Similarity 100.0%; Pred. NO. 0.0087;  
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
Qy 1 CCTTAGGAGGAACAAGTCCC 20  
Db 2899 CCTTAGGAGGAACAAGTCCC 2918  
RESULT 6  
US-09-620-312D-823  
Sequence 823, Application US/09620312D  
Patent No. 6569662  
GENERAL INFORMATION:  
APPLICANT: Tang, Y. Tom  
APPLICANT: Liu, Chenghua  
APPLICANT: Asundi, Vinod  
APPLICANT: Zhang, Jie

APPLICANT: Ren, Feiyan  
APPLICANT: Chen, Rui-hong  
APPLICANT: Zhao, Qing A.  
APPLICANT: Wehrman, Tom  
APPLICANT: Xue, Aidong J.  
APPLICANT: Yang, Yonghong  
APPLICANT: Wang, Jian-Rui  
APPLICANT: Zhou, Ping  
APPLICANT: Ma, Yungding  
APPLICANT: Wang, Dunrui  
APPLICANT: Wang, Zhiwei  
APPLICANT: John Tillinghast  
APPLICANT: Drmanac, Radojac T.  
TITLE OF INVENTION: No.6569662e1 Nucleic Acids and  
TITLE OF INVENTION: Polypeptides  
FILE REFERENCE: 784C1P2B  
CURRENT APPLICATION NUMBER: US/09/620,312D  
CURRENT FILING DATE: 2000-07-19  
PRIOR APPLICATION NUMBER: 09/552,317  
PRIOR FILING DATE: 2000-04-25  
PRIOR APPLICATION NUMBER: 09/488,725  
PRIOR FILING DATE: 2000-01-21  
NUMBER OF SEQ ID NOS: 1105  
SOFTWARE: pc FL\_genes Version 1.0  
SEQ ID NO 823  
LENGTH: 6572  
TYPE: DNA  
ORGANISM: Homo sapiens  
FEATURE:  
NAME/KEY: CDS  
LOCATION: (521)..(2611)  
US-09-620-312D-823

Query Match 80.0%; Score 16; DB 4; Length 6572;  
Best Local Similarity 100.0%; Pred. No. 1.4;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 CTTAGAGGAAACAGT 17  
DB 4524 CTTAGAGGAAACAGT 4539

RESULT 7  
US-09-439-313-354  
Sequence 354, Application US/09439313  
Patent No. 6329505  
GENERAL INFORMATION:  
APPLICANT: Xu, Jiangchun  
APPLICANT: Dillon, Davin C.  
APPLICANT: Mitchem, Jennifer L.  
APPLICANT: Harlocker, Susan Louise  
APPLICANT: Jiang Yuqi  
APPLICANT: Reed, Steven G.  
APPLICANT: Kalos, Michael  
APPLICANT: Fanger, Gary  
APPLICANT: Retter, Mark  
APPLICANT: Solk, John  
APPLICANT: Day, Craig  
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THERAPY AND  
TITLE OF INVENTION: DIAGNOSIS OF PROSTATE CANCER  
FILE REFERENCE: 210121.427C9  
CURRENT APPLICATION NUMBER: US/09/439,313  
CURRENT FILING DATE: 1999-11-12  
NUMBER OF SEQ ID NOS: 575  
SOFTWARE: FastSeq for Windows Version 3.0  
SEQ ID NO 354  
LENGTH: 854  
TYPE: DNA  
ORGANISM: Homo sapien  
US-09-439-313-354  
Query Match 70.0%; Score 14; DB 4; Length 854;  
Best Local Similarity 100.0%; Pred. No. 20;

Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 5 AGGAGGAACAGTC 18  
DB 497 AGGAGGAACAGTC 510

RESULT 8  
US-09-352-616A-354  
Sequence 354, Application US/09352616A  
Patent No. 6395278  
GENERAL INFORMATION:  
APPLICANT: Dillon, Davin C.  
APPLICANT: Harlocker, Susan Louise  
APPLICANT: Jiang Yuqi  
APPLICANT: Xu, Jiangchun  
APPLICANT: Mitchem, Jennifer Lynn  
TITLE OF INVENTION: COMPOUNDS FOR IMMUNOTHERAPY AND DIAGNOSIS  
TITLE OF INVENTION: OF PROSTATE CANCER AND METHODS FOR THEIR USE  
FILE REFERENCE: 210121.427C8  
CURRENT APPLICATION NUMBER: US/09/352,616A  
CURRENT FILING DATE: 1999-07-13  
NUMBER OF SEQ ID NOS: 472  
SOFTWARE: FastSeq for Windows Version 3.0  
SEQ ID NO 354  
LENGTH: 854  
TYPE: DNA  
ORGANISM: Homo sapien  
US-09-352-616A-354

Query Match 70.0%; Score 14; DB 4; Length 854;  
Best Local Similarity 100.0%; Pred. No. 20;  
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 AGGAGGAACAGTC 18  
DB 497 AGGAGGAACAGTC 510

RESULT 9  
US-08-599-171A-17/c  
Sequence 17, Application US/08599171A  
Patent No. 5814473  
GENERAL INFORMATION:  
APPLICANT: WARREN, Patrick V.  
TITLE OF INVENTION: TRANSAMINASES AND AMINOTRANSFERASES  
NUMBER OF SEQUENCES: 32  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: CARELLA, BYRNE, BAIN, GILFILLAN,  
ADDRESSEE: CECCHI, STEWART & OLSTEIN  
STREET: 6 BECKER FARM ROAD  
CITY: ROSELAND  
STATE: NEW JERSEY  
COUNTRY: USA  
ZIP: 07068  
COMPUTER READABLE FORM:  
MEDIUM TYPE: 3.5 INCH DISKETTE  
COMPUTER: IBM PS/2  
OPERATING SYSTEM: MS-DOS  
SOFTWARE: WORD PERFECT 5.1  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/599,171A  
FILING DATE: Concurrently  
CLASSIFICATION: 435  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER:  
FILING DATE:  
CLASSIFICATION: 435  
ATTORNEY/AGENT INFORMATION:  
NAME: HERON, CHARLES J.  
REGISTRATION NUMBER: 28,019  
REFERENCE/DOCKET NUMBER: 331400-38  
TELECOMMUNICATION INFORMATION:

TELEPHONE: 201-994-1700  
TELEFAX: 201-994-1744  
INFORMATION FOR SEQ ID NO: 17:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 1245 NUCLEOTIDES  
TYPE: NUCLEIC ACID  
STRANDEDNESS: SINGLE  
TOPOLOGY: LINEAR  
MOLECULE TYPE: GENOMIC DNA  
US-08-599-171A-17

Query Match 70.0%; Score 14; DB 1; Length 1245;  
Best Local Similarity 100.0%; Pred. No. 19;  
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 3 TTAGAGGAACAAG 16  
|||||  
Db 1069 TTAGAGGAACAAG 1056

RESULT 10  
US-08-646-590B-17/C  
Sequence 17, Application US/08646590B  
Patent No. 5962283  
GENERAL INFORMATION:  
APPLICANT: Warren, Patrick V.  
APPLICANT: Swanson, Ronald V.  
TITLE OF INVENTION: TRANSAMINASES AND AMINOTRANSFERASES  
NUMBER OF SEQUENCES: 42  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Fish & Richardson, P.C.  
STREET: 4225 Executive Square, Suite 1400  
CITY: La Jolla  
STATE: CA  
COUNTRY: US  
ZIP: 92037  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Diskette  
COMPUTER: IBM Compatible  
OPERATING SYSTEM: Windows95  
SOFTWARE: FASTSEQ for Windows Version 2.0  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/646,590B  
FILING DATE: 08-May-1996  
CLASSIFICATION: 435  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 08/599,171  
FILING DATE: 09-FEB-1996  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: PCT/US97/01094  
FILING DATE: 21-January-1997  
CLASSIFICATION: 435  
ATTORNEY/AGENT INFORMATION:  
NAME: Halle, Ph.D., Lisa A.  
REGISTRATION NUMBER: 38,347  
REFERENCE/DOCKET NUMBER: 09010/017001  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 619/678-5070  
TELEFAX: 619/678-5099  
INFORMATION FOR SEQ ID NO: 17:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 1245 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
MOLECULE TYPE: Genomic DNA  
FEATURE:  
NAME/KEY: Coding Sequence  
LOCATION: 1...1242  
US-08-646-590B-17

Query Match 70.0%; Score 14; DB 2; Length 1245;  
Best Local Similarity 100.0%; Pred. No. 19;

Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
OY 3 TTAGAGGAACAAG 16  
|||||  
Db 1069 TTAGAGGAACAAG 1056

RESULT 11  
US-09-069-226-17/C  
Sequence 17, Application US/09069226  
Patent No. 6013509  
GENERAL INFORMATION:  
APPLICANT: WARREN, Patrick V.  
TITLE OF INVENTION: TRANSAMINASES AND AMINOTRANSFERASES  
NUMBER OF SEQUENCES: 32  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: CARELLA, BYRNE, BAIN, GILFILLAN,  
ADDRESSEE: CECCHI, STEWART & OLSTEIN  
STREET: 6 BECKER FARM ROAD  
CITY: ROSELAND  
STATE: NEW JERSEY  
COUNTRY: USA  
ZIP: 07068  
COMPUTER READABLE FORM:  
MEDIUM TYPE: 3.5 INCH DISKETTE  
COMPUTER: IBM PS/2  
OPERATING SYSTEM: MS-DOS  
SOFTWARE: WORD PERFECT 5.1  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/09/069,226  
FILING DATE:  
CLASSIFICATION:  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 08/599,171  
FILING DATE:  
ATTORNEY/AGENT INFORMATION:  
NAME: HERRON, CHARLES J.  
REGISTRATION NUMBER: 28,019  
REFERENCE/DOCKET NUMBER: 331400-38  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 201-994-1700  
TELEFAX: 201-994-1744  
INFORMATION FOR SEQ ID NO: 17:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 1245 NUCLEOTIDES  
TYPE: NUCLEIC ACID  
STRANDEDNESS: SINGLE  
TOPOLOGY: LINEAR  
MOLECULE TYPE: GENOMIC DNA  
US-09-069-226-17

Query Match 70.0%; Score 14; DB 3; Length 1245;  
Best Local Similarity 100.0%; Pred. No. 19;  
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 3 TTAGAGGAACAAG 16  
|||||  
Db 1069 TTAGAGGAACAAG 1056

RESULT 12  
US-09-412-184-17/C  
Sequence 17, Application US/09412184  
Patent No. 6268188  
GENERAL INFORMATION:  
APPLICANT: Warren, Patrick V.  
APPLICANT: Swanson, Ronald V.  
TITLE OF INVENTION: TRANSAMINASES AND AMINOTRANSFERASES  
NUMBER OF SEQUENCES: 42  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Fish & Richardson, P.C.  
STREET: 4225 Executive Square, Suite 1400  
CITY: La Jolla

STATE: CA  
COUNTRY: US  
Z12: 92037  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Diskette  
COMPUTER: IBM Compatible  
OPERATING SYSTEM: Windows95  
SOFTWARE: FastSeq for Windows Version 2.0  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/09/412.184  
FILING DATE:  
CLASSIFICATION:  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US/08/646.590  
FILING DATE: 08-May-1996  
APPLICATION NUMBER: 08/599.171  
FILING DATE: 09-FEB-1996  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: PCT/US97/01094  
FILING DATE: 21-January-1997  
CLASSIFICATION:  
ATTORNEY/AGENT INFORMATION:  
NAME: Haile, Ph.D., Lisa A.  
REGISTRATION NUMBER: 38,347  
REFERENCE/DOCKET NUMBER: 09010/017001  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 619/678-5070  
TELEFAX: 619/678-5099  
INFORMATION FOR SEQ ID NO: 17:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 1245 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
MOLECULE TYPE: Genomic DNA  
FEATURE:  
NAME/KEY: Coding Sequence  
LOCATION: 1...1242  
US-09-412-184-17

Query Match 70.0%; Score 14; DB 3; Length 1245;  
Best Local Similarity 100.0%; Pred. No. 19;  
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 TTAGGAGGACAG 16  
DB 1069 TTAGGAGGACAG 1056

RESULT 13  
US-09-232-160-6/C  
Sequence 6, Application US/09222160  
Patent No. 6368794  
GENERAL INFORMATION:  
APPLICANT: Steve Daniel  
APPLICANT: James Gilmore  
APPLICANT: Susan G. Stuart  
APPLICANT: Laura Stuve  
TITLE OF INVENTION: DETECTION OF ALTERED EXPRESSION OF GENES REGULATING CELL  
FILE REFERENCE: PA-0003 US  
CURRENT APPLICATION NUMBER: US/09/232,160  
FILING DATE: 1999-01-15  
NUMBER OF SEQ ID NOS: 23  
SOFTWARE: PERL Program  
SEQ ID NO 6  
LENGTH: 1585  
TYPE: DNA  
ORGANISM: Homo sapiens  
FEATURE:  
OTHER INFORMATION: 1698542  
US-09-232-160-6

Query Match 70.0%; Score 14; DB 4; Length 1585;  
Best Local Similarity 100.0%; Pred. No. 19;  
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 7 GAGGACAGTCCC 20  
DB 536 GAGGACAGTCCC 523

RESULT 14  
US-08-460-784-21/C  
Sequence 21, Application US/08460784  
Patent No. 5693473  
GENERAL INFORMATION:  
APPLICANT: Skolnick, Mark H.  
APPLICANT: Goldgar, David E.  
APPLICANT: Mikl, Yoshio  
APPLICANT: Swenson, Jeff  
APPLICANT: Kamb, Alexander  
APPLICANT: Harshman, Keith D.  
APPLICANT: Shattuck-Bidens, Donna M.  
APPLICANT: Tavtigian, Sean V.  
APPLICANT: Wiseman, Roger W.  
TITLE OF INVENTION: 17q-Linked Breast and Ovarian Cancer  
NUMBER OF SEQUENCES: 85  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Venable, Baetjer, Howard & Civiletti, LLP  
STREET: 1201 New York Avenue, N.W., Suite 1000  
CITY: Washington  
STATE: DC  
COUNTRY: USA  
ZIP: 20005

COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: Patent Release #1.0, Version #1.30  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/460,784  
FILING DATE:  
CLASSIFICATION: 435  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/409,305  
FILING DATE: 24-MAR-1995  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/348,824  
FILING DATE: 29-NOV-1994  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/308,104  
FILING DATE: 16-SEP-1994  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/300,266  
FILING DATE: 02-SEP-1994  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/289,221  
FILING DATE: 12-AUG-1994  
ATTORNEY/AGENT INFORMATION:  
NAME: Ihnen, Jeffrey L.  
REGISTRATION NUMBER: 28,957  
REFERENCE/DOCKET NUMBER: 24884-109347  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 202-962-4810  
TELEFAX: 202-962-8300  
INFORMATION FOR SEQ ID NO: 21:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 4249 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: double  
TOPOLOGY: linear  
MOLECULE TYPE: DNA (genomic)  
HYPOTHETICAL: NO

ANTI-SENSE: NO  
ORIGINAL SOURCE:  
ORGANISM: Homo sapiens  
US-08-480-784-21

Query Match 70.0%; Score 14; DB 1; Length 4249;  
Best Local Similarity 100.0%; Pred. No. 18;  
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 CCTAGAGAGACA 14  
DB 3770 CCTAGAGAGACA 3757

RESULT 15  
US-08-483-553-21/c  
Sequence 21, Application US/08483553  
Patent No. 5709999  
GENERAL INFORMATION:  
APPLICANT: Skolnick, Mark H.  
APPLICANT: Goldgar, David E.  
APPLICANT: Miki, Yoshio  
APPLICANT: Swenson, Jeff  
APPLICANT: Kamd, Alexander  
APPLICANT: Hartsman, Keith D.  
APPLICANT: Shattuck-Eidens, Donna M.  
APPLICANT: Tavligian, Sean V.  
APPLICANT: Wiseman, Roger W.  
APPLICANT: Futreal, P. Andrew  
TITLE OF INVENTION: 17q-Linked Breast and Ovarian Cancer  
TITLE OF INVENTION: Susceptibility Gene  
NUMBER OF SEQUENCES: 85  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Venable, Baetjer, Howard & Civiletti, LLP  
STREET: 1201 New York Avenue, N.W., Suite 1000  
CITY: Washington  
STATE: DC  
COUNTRY: USA  
ZIP: 20005  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: Patentin Release #1.0, Version #1.30  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/483,553  
FILING DATE:  
CLASSIFICATION: 435  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/409,305  
FILING DATE: 24-MAR-1995  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/348,824  
FILING DATE: 29-NOV-1994  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/308,104  
FILING DATE: 16-SEP-1994  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/300,266  
FILING DATE: 02-SEP-1994  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/289,221  
FILING DATE: 12-AUG-1994  
ATTORNEY/AGENT INFORMATION:  
NAME: Ihnen, Jeffrey L.  
REGISTRATION NUMBER: 28,957  
REFERENCE/DOCKET NUMBER: 24884-109347  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 202-962-4810  
TELEFAX: 202-962-8300  
INFORMATION FOR SEQ ID NO: 21:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 4249 base pairs

TYPE: nucleic acid  
STRANDEDNESS: double  
TOPOLOGY: linear  
MOLECULE TYPE: DNA (genomic)  
HYPOTHETICAL: NO  
ANTI-SENSE: NO  
ORIGINAL SOURCE:  
ORGANISM: Homo sapiens  
US-08-483-553-21

Query Match 70.0%; Score 14; DB 1; Length 4249;  
Best Local Similarity 100.0%; Pred. No. 18;  
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 CCTAGAGAGACA 14  
DB 3770 CCTAGAGAGACA 3757

RESULT 16  
US-08-487-002-21/c  
Sequence 21, Application US/08487002  
Patent No. 5710001  
GENERAL INFORMATION:  
APPLICANT: Shattuck-Eidens, Donna M.  
APPLICANT: Simard, Jacques  
APPLICANT: Eml, Mitsuru  
APPLICANT: Nakamura, Yusuke  
APPLICANT: Durocher, Francine  
TITLE OF INVENTION: 17q-Linked Breast and Ovarian Cancer  
TITLE OF INVENTION: Susceptibility Gene  
NUMBER OF SEQUENCES: 85  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Venable, Baetjer, Howard & Civiletti, LLP  
STREET: 1201 New York Avenue, N.W., Suite 1000  
CITY: Washington  
STATE: DC  
COUNTRY: USA  
ZIP: 20005  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: Patentin Release #1.0, Version #1.30  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/487,002  
FILING DATE:  
CLASSIFICATION: 424  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/409,305  
FILING DATE: 24-MAR-1995  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/348,824  
FILING DATE: 29-NOV-1994  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/308,104  
FILING DATE: 16-SEP-1994  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/300,266  
FILING DATE: 02-SEP-1994  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/289,221  
FILING DATE: 12-AUG-1994  
ATTORNEY/AGENT INFORMATION:  
NAME: Ihnen, Jeffrey L.  
REGISTRATION NUMBER: 28,957  
REFERENCE/DOCKET NUMBER: 24884-109347  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 202-962-4810  
TELEFAX: 202-962-8300  
INFORMATION FOR SEQ ID NO: 21:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 4249 base pairs

TYPE: nucleic acid.  
STRANDEDNESS: double  
TOPOLOGY: linear  
MOLECULE TYPE: DNA (genomic)  
HYPOTHEICAL: NO  
ANTI-SENSE: NO  
ORIGINAL SOURCE:  
ORGANISM: Homo sapiens  
US-08-487-002-21

Query Match 70.0%; Score 14; DB 1; Length 4249;  
Best Local Similarity 100.0%; Pred. No. 18;  
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CCTTAGGAGGACCA 14  
Db 3770 CCTTAGGAGGACCA 3757

RESULT 17  
US-08-483-554B-21/C  
Sequence 21, Application US/08483554B  
Patent No. 5747282  
GENERAL INFORMATION:  
APPLICANT: Skolnick, Mark H.  
APPLICANT: Goldgar, David E.  
APPLICANT: Miki, Yoshio  
APPLICANT: Swenson, Jeff  
APPLICANT: Kamb, Alexander  
APPLICANT: Harshman, Keith D.  
APPLICANT: Shattuck-Eidens, Donna M.  
APPLICANT: Tavligian, Sean V.  
APPLICANT: Wiseman, Roger W.  
APPLICANT: Futreal, P. Andrew  
TITLE OF INVENTION: 17q-Linked Breast and Ovarian Cancer  
NUMBER OF SEQUENCES: 85  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Venable, Baetjer, Howard & Civiletti, LLP  
STREET: 1201 New York Avenue, N.W., Suite 1000  
CITY: Washington  
STATE: DC  
COUNTRY: USA  
ZIP: 20005  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: Patentin Release #1.0, Version #1.30  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/483,554B  
FILING DATE: 07-JUN-1995  
CLASSIFICATION: 514  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/409,305  
FILING DATE: 24-MAR-1995  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/348,824  
FILING DATE: 29-NOV-1994  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/308,104  
FILING DATE: 16-SEP-1994  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/300,266  
FILING DATE: 02-SEP-1994  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/289,221  
FILING DATE: 12-AUG-1994  
ATTORNEY/AGENT INFORMATION:  
NAME: Ihnen, Jeffrey L.  
REGISTRATION NUMBER: 28,957  
REFERENCE/DOCKET NUMBER: 24884-109347  
TELECOMMUNICATION INFORMATION:

TELEPHONE: 202-962-4810  
TELEFAX: 202-962-8300  
INFORMATION FOR SEQ ID NO: 21:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 4249 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: double  
TOPOLOGY: linear  
MOLECULE TYPE: DNA (genomic)  
HYPOTHEICAL: NO  
ANTI-SENSE: NO  
ORIGINAL SOURCE:  
ORGANISM: Homo sapiens  
US-08-483-554B-21

Query Match 70.0%; Score 14; DB 1; Length 4249;  
Best Local Similarity 100.0%; Pred. No. 18;  
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CCTTAGGAGGACCA 14  
Db 3770 CCTTAGGAGGACCA 3757

RESULT 18  
US-08-488-011B-21/C  
Sequence 21, Application US/08488011B  
Patent No. 5751441  
GENERAL INFORMATION:  
APPLICANT: Skolnick, Mark H.  
APPLICANT: Goldgar, David E.  
APPLICANT: Miki, Yoshio  
APPLICANT: Swenson, Jeff  
APPLICANT: Kamb, Alexander  
APPLICANT: Harshman, Keith D.  
APPLICANT: Shattuck-Eidens, Donna M.  
APPLICANT: Tavligian, Sean V.  
APPLICANT: Wiseman, Roger W.  
APPLICANT: Futreal, P. Andrew  
TITLE OF INVENTION: 17q-Linked Breast and Ovarian Cancer  
NUMBER OF SEQUENCES: 85  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Venable, Baetjer, Howard & Civiletti, LLP  
STREET: 1201 New York Avenue, N.W., Suite 1000  
CITY: Washington  
STATE: DC  
COUNTRY: USA  
ZIP: 20005  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: Patentin Release #1.0, Version #1.30  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/488,011B  
FILING DATE: 07-JUN-1995  
CLASSIFICATION: 435  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/409,305  
FILING DATE: 24-MAR-1995  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/348,824  
FILING DATE: 29-NOV-1994  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/308,104  
FILING DATE: 16-SEP-1994  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/300,266  
FILING DATE: 02-SEP-1994  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/289,221  
FILING DATE: 12-AUG-1994



ATTORNEY/AGENT INFORMATION:  
NAME: Ihnen, Jeffrey L.  
REGISTRATION NUMBER: 28,957  
REFERENCE/DOCKET NUMBER: 24884-109347-09  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 202-962-4810  
TELEFAX: 202-962-8300  
INFORMATION FOR SEQ ID NO: 21:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 4249 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: double  
TOPOLOGY: linear  
MOLECULE TYPE: DNA (genomic)  
HYPOTHETICAL: NO  
ANTI-SENSE: NO  
ORIGINAL SOURCE:  
ORGANISM: Homo sapiens  
US-08-488-011B-21

Query Match 70.0%; Score 14; DB 1; Length 4249;  
Best Local Similarity 100.0%; Pred. No. 18;  
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CCTTAGGAGGACA 14  
Db 3770 CCTTAGGAGGACA 3757

RESULT 19  
US-08-850-727-21/c  
Sequence 21, Application US/08850727  
Patent No. 6162897  
GENERAL INFORMATION:  
APPLICANT: Skolnick, Mark H.  
APPLICANT: Goldgar, David E.  
APPLICANT: Miki, Yoshio  
APPLICANT: Swenson, Jeff  
APPLICANT: Kamb, Alexander  
APPLICANT: Harshman, Keith D.  
APPLICANT: Shattuck-Eidens, Donna M.  
APPLICANT: Tavligian, Sean V.  
APPLICANT: Wiseman, Roger W.  
APPLICANT: Futreal, P. Andrew  
TITLE OF INVENTION: 17q-linked Breast and Ovarian Cancer  
TITLE OF INVENTION: Susceptibility Gene  
NUMBER OF SEQUENCES: 85  
CORRESPONDENCE ADDRESSES:  
ADDRESSEE: Venable, Baetjer, Howard & Civiletti, LLP  
STREET: 1201 New York Avenue, N.W., Suite 1000  
CITY: Washington  
STATE: DC  
COUNTRY: USA  
ZIP: 20005  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: Patentin Release #1.0, Version #1.30  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/850,727  
FILING DATE:  
CLASSIFICATION:  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/483,554  
FILING DATE: 07-JUN-1995  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/348,824  
FILING DATE: 29-NOV-1994  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/308,104  
FILING DATE: 16-SEP-1994  
PRIOR APPLICATION DATA:

APPLICATION NUMBER: US 08/300,266  
FILING DATE: 02-SEP-1994  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/289,221  
FILING DATE: 12-AUG-1994  
ATTORNEY/AGENT INFORMATION:  
NAME: Ihnen, Jeffrey L.  
REGISTRATION NUMBER: 28,957  
REFERENCE/DOCKET NUMBER: 24884-109347  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 202-962-4810  
TELEFAX: 202-962-8300  
INFORMATION FOR SEQ ID NO: 21:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 4249 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: double  
TOPOLOGY: linear  
MOLECULE TYPE: DNA (genomic)  
HYPOTHETICAL: NO  
ANTI-SENSE: NO  
ORIGINAL SOURCE:  
ORGANISM: Homo sapiens  
US-08-850-727-21

Query Match 70.0%; Score 14; DB 3; Length 4249;  
Best Local Similarity 100.0%; Pred. No. 18;  
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CCTTAGGAGGACA 14  
Db 3770 CCTTAGGAGGACA 3757

RESULT 20  
PCT-US95-10202-21/c  
Sequence 21, Application PC/TUS9510202  
GENERAL INFORMATION:  
APPLICANT: Shattuck-Eidens, Donna M.  
APPLICANT: Simard, Jacques  
APPLICANT: Emi, Mitsuru  
APPLICANT: Nakamura, Yusuke  
APPLICANT: Durocher, Francine  
TITLE OF INVENTION: In Vivo Mutations and Polymorphisms  
TITLE OF INVENTION: in the 17q-linked Breast and Ovarian Cancer  
TITLE OF INVENTION: Susceptibility Gene  
NUMBER OF SEQUENCES: 85  
CORRESPONDENCE ADDRESSES:  
ADDRESSEE: Venable, Baetjer, Howard & Civiletti, LLP  
STREET: 1201 New York Avenue, N.W., Suite 1000  
CITY: Washington  
STATE: DC  
COUNTRY: USA  
ZIP: 20005  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: Patentin Release #1.0, Version #1.30  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: PCT/US95/10202  
FILING DATE:  
CLASSIFICATION:  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US  
FILING DATE: 07-JUN-1995  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/409,305  
FILING DATE: 24-MAR-1995  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/348,824  
FILING DATE: 29-NOV-1994  
PRIOR APPLICATION DATA:

APPLICATION NUMBER: US 08-308,104  
FILING DATE: 16-SEP-1994  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/300,266  
FILING DATE: 02-SEP-1994  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/289,221  
FILING DATE: 12-AUG-1994  
ATTORNEY/AGENT INFORMATION:  
NAME: Ihnen, Jeffrey L.  
REGISTRATION NUMBER: 28,957  
REFERENCE/DOCKET NUMBER: 24884-109347  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 202-962-8300  
TELEFAX: 202-962-8300  
INFORMATION FOR SEQ ID NO: 21:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 4249 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: double  
TOPOLOGY: linear  
MOLECULE TYPE: DNA (genomic)  
HYPOTHETICAL: NO  
ANTI-SENSE: NO  
ORIGINAL SOURCE:  
ORGANISM: Homo sapiens  
PCT-US95-10202-21

Query Match 70.0%; Score 14; DB 5; Length 4249;  
Best Local Similarity 100.0%; Pred. No. 18;  
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 CCTAGAGAGACA 14  
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Db 3770 CCTAGAGAGACA 3757

RESULT 21  
PCT-US95-10203-21/c  
Sequence 21, Application PC/TUS9510203  
GENERAL INFORMATION:  
APPLICANT: Skolnick, Mark H.  
APPLICANT: Goldgar, David E.  
APPLICANT: Miki, Yoshio  
APPLICANT: Swenson, Jeff  
APPLICANT: Kamb, Alexander  
APPLICANT: Hershman, Keith D.  
APPLICANT: Shattuck-Eidens, Donna M.  
APPLICANT: Tavtigian, Sean V.  
APPLICANT: Wiseman, Roger W.  
APPLICANT: Futreal, P. Andrew  
TITLE OF INVENTION: 17q-linked Breast and Ovarian Cancer  
TITLE OF INVENTION: Susceptibility Gene  
NUMBER OF SEQUENCES: 85  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Venable, Baetjer, Howard & Civiletti, LLP  
STREET: 1201 New York Avenue, N.W., Suite 1000  
CITY: Washington  
STATE: DC  
COUNTRY: USA  
ZIP: 20005  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: Patent Release #1.0, Version #1.30  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: PCT/US95/10203  
FILING DATE:  
CLASSIFICATION:  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US  
FILING DATE: 07-JUN-1995

PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/409,305  
FILING DATE: 24-MAR-1995  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/348,824  
FILING DATE: 29-NOV-1994  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08-308,104  
FILING DATE: 16-SEP-1994  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/300,266  
FILING DATE: 02-SEP-1994  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/289,221  
FILING DATE: 12-AUG-1994  
ATTORNEY/AGENT INFORMATION:  
NAME: Ihnen, Jeffrey L.  
REGISTRATION NUMBER: 28,957  
REFERENCE/DOCKET NUMBER: 24884-109347  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 202-962-8300  
TELEFAX: 202-962-8300  
INFORMATION FOR SEQ ID NO: 21:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 4249 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: double  
TOPOLOGY: linear  
MOLECULE TYPE: DNA (genomic)  
HYPOTHETICAL: NO  
ANTI-SENSE: NO  
ORIGINAL SOURCE:  
ORGANISM: Homo sapiens  
PCT-US95-10203-21

Query Match 70.0%; Score 14; DB 5; Length 4249;  
Best Local Similarity 100.0%; Pred. No. 18;  
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 CCTAGAGAGACA 14  
|||  
Db 3770 CCTAGAGAGACA 3757

RESULT 22  
PCT-US95-10220-21/c  
Sequence 21, Application PC/TUS9510220  
GENERAL INFORMATION:  
APPLICANT: Skolnick, Mark H.  
APPLICANT: Goldgar, David E.  
APPLICANT: Miki, Yoshio  
APPLICANT: Swenson, Jeff  
APPLICANT: Kamb, Alexander  
APPLICANT: Hershman, Keith D.  
APPLICANT: Shattuck-Eidens, Donna M.  
APPLICANT: Tavtigian, Sean V.  
APPLICANT: Wiseman, Roger W.  
APPLICANT: Futreal, P. Andrew  
TITLE OF INVENTION: Method for Diagnosing a  
TITLE OF INVENTION: Predisposition for Breast and Ovarian Cancer  
NUMBER OF SEQUENCES: 85  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Venable, Baetjer, Howard & Civiletti, LLP  
STREET: 1201 New York Avenue, N.W., Suite 1000  
CITY: Washington  
STATE: DC  
COUNTRY: USA  
ZIP: 20005  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: Patent Release #1.0, Version #1.30

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; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: PCT/US95/10220
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US
; FILING DATE: 07-JUN-1995
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/409,305
; FILING DATE: 24-MAR-1995
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/348,824
; FILING DATE: 29-NOV-1994
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08-308,104
; FILING DATE: 16-SEP-1994
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/300,266
; FILING DATE: 02-SEP-1994
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/289,221
; FILING DATE: 12-AUG-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: Immen, Jeffrey L.
; REGISTRATION NUMBER: 28,957
; REFERENCE/DOCKET NUMBER: 24884-109347
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 202-962-4810
; TELEFAX: 202-962-8300
; INFORMATION FOR SEQ ID NO: 21:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 4249 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; HYPOTHEetical: NO
; ANTI-SENSE: NO
; ORIGINAL SOURCE:
; ORGANISM: Homo sapiens
; PCT-US95-10220-21

Query March 70.0%; Score 14; DB 5; Length 4249;
Best Local Similarity 100.0%; Pred. No. 18;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CCTTAGAGAGACA 14
Db 3770 CCTTAGAGAGACA 3757

RESULT 23
US-08-746-411A-9/c
; Sequence 9, Application US/08746411A
; Patent No. 6117632
; GENERAL INFORMATION:
; APPLICANT: O'Mahony, Daniel J
; TITLE OF INVENTION: Peptides Which Enhance Transport Across
; TITLE OF INVENTION: Tissues and Methods of Identifying and Using the Same
; NUMBER OF SEQUENCES: 17
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Mary L. Severson, Ph.D., Esq.
; STREET: 1300 Gould Drive
; CITY: Gainesville
; STATE: GA
; COUNTRY: USA
; ZIP: 30504
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:

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; APPLICATION NUMBER: US/08/746,411A
; FILING DATE: 08-NOV-1996
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 60/006461
; FILING DATE: 10-NOV-1995
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: IE 950865
; FILING DATE: 10-NOV-1995
; ATTORNEY/AGENT INFORMATION:
; NAME: Severson, Mary L.
; REGISTRATION NUMBER: 34,927
; REFERENCE/DOCKET NUMBER: 96.1060. US
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 770 534-8239
; TELEFAX: 770 534-8247
; INFORMATION FOR SEQ ID NO: 9:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 126 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: other nucleic acid
; DESCRIPTION: /desc = "individual isolate"
; US-08-746-411A-9

Query March 65.0%; Score 13; DB 3; Length 126;
Best Local Similarity 100.0%; Pred. No. 79;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 TAGGAGAGACAG 16
Db 37 TAGGAGAGACAG 25

RESULT 24
US-08-857-046A-9/c
; Sequence 9, Application US/08857046A
; Patent No. 6361938
; GENERAL INFORMATION:
; APPLICANT: O'Mahony, Daniel J
; APPLICANT: Alvarez, Vernon L
; APPLICANT: Seveso, Michele
; TITLE OF INVENTION: Peptides Which Enhance Transport Across
; TITLE OF INVENTION: Tissues and Methods of Identifying and Using the Same
; NUMBER OF SEQUENCES: 33
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Mary L. Severson, Ph.D., Esq.
; STREET: 1300 Gould Drive
; CITY: Gainesville
; STATE: GA
; COUNTRY: USA
; ZIP: 30504
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/857,046A
; FILING DATE: 15-MAY-1997
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 60/006461
; FILING DATE: 10-NOV-1995
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: IE 950864
; FILING DATE: 10-NOV-1995
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/746,411
; FILING DATE: 08-NOV-1996
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: WO PCT/IE96/00073

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FILING DATE: 11-NOV-1996  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: WO PCT/IE96/00072  
FILING DATE: 11-NOV-1996  
ATTORNEY/AGENT INFORMATION:  
NAME: Severson, Mary L  
REGISTRATION NUMBER: 34,927  
REFERENCE/DOCKET NUMBER: 97,1061.US  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 770 534-8239  
TELEFAX: 770 534-8247  
INFORMATION FOR SEQ ID NO: 9:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 126 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
MOLECULE TYPE: other nucleic acid  
DESCRIPTION: /desc = "individual isolate"  
US-08-857-046A-9

Query Match 65.0%; Score 13; DB 4; Length 126;  
Best Local Similarity 100.0%; Pred. No. 79;  
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 4 TAGAGGAGAACAG 16  
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Db 37 TAGAGGAGAACAG 25

RESULT 25  
US-09-573-252-9/c  
Sequence 9, Application US/09573252  
Patent No. 6521737  
GENERAL INFORMATION:  
APPLICANT: O'Mahony, Daniel J  
TITLE OF INVENTION: Peptides Which Enhance Transport Across  
Tissues and Methods of Identifying and Using the Same  
NUMBER OF SEQUENCES: 17  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Mary L. Severson, Ph.D., Esq.  
STREET: 1300 Gould Drive  
CITY: Gainesville  
STATE: GA  
COUNTRY: USA  
ZIP: 30604  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: Patentin Release #1.0, Version #1.30  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/09/573,252  
FILING DATE: 19-Aug-2002  
CLASSIFICATION: <Unknown>  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US/08/746,411  
FILING DATE: 08-NOV-1996  
APPLICATION NUMBER: US 60/006461  
FILING DATE: 10-NOV-1995  
APPLICATION NUMBER: IE 950865  
FILING DATE: 10-NOV-1995  
ATTORNEY/AGENT INFORMATION:  
NAME: Severson, Mary L  
REGISTRATION NUMBER: 34,927  
REFERENCE/DOCKET NUMBER: 96,1060.US  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 770 534-8239  
TELEFAX: 770 534-8247  
INFORMATION FOR SEQ ID NO: 9:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 126 base pairs  
TYPE: nucleic acid

STRANDEDNESS: single  
TOPOLOGY: linear  
MOLECULE TYPE: other nucleic acid  
DESCRIPTION: /desc = "individual isolate"  
US-09-573-252-9

Query Match 65.0%; Score 13; DB 4; Length 126;  
Best Local Similarity 100.0%; Pred. No. 79;  
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 4 TAGAGGAGAACAG 16  
|||  
Db 37 TAGAGGAGAACAG 25

RESULT 26  
US-08-248-474-23/c  
Sequence 23, Application US/08248474  
Patent No. 5612471  
GENERAL INFORMATION:  
APPLICANT: MCK, BIRD, David  
TITLE OF INVENTION: NEMATODE-INDUCE GENES IN TOMATO  
NUMBER OF SEQUENCES: 114  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Townsend and Townsend Kourie and Crew  
STREET: Steuart Street Tower, One Market Plaza  
CITY: San Francisco  
STATE: California  
COUNTRY: US  
ZIP: 94105-1493  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: Patentin Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/248,474  
FILING DATE: 25-MAY-1994  
CLASSIFICATION: 800  
ATTORNEY/AGENT INFORMATION:  
NAME: Bastian, Kevin L.  
REGISTRATION NUMBER: 34,774  
REFERENCE/DOCKET NUMBER: 2307E-535  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: (415) 543-9600  
TELEFAX: (415) 543-5043  
INFORMATION FOR SEQ ID NO: 23:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 243 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
MOLECULE TYPE: DNA (genomic)  
ORIGINAL SOURCE:  
ORGANISM: Lycopersicon esculentum cv 'Rutgers large  
ORGANISM: Red'  
FEATURE:  
NAME/KEY: misc\_feature  
LOCATION: 1..243  
OTHER INFORMATION: /standard\_name= "DB# 139"  
US-08-248-474-23

Query Match 65.0%; Score 13; DB 1; Length 243;  
Best Local Similarity 100.0%; Pred. No. 76;  
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 3 TTAGAGGAGAACAA 15  
|||  
Db 133 TTAGAGGAGAACAA 121

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RESULT 27
US-08-756-849-23/c
; Sequence 23, Application US/08756849
; Patent No. 6093810
; GENERAL INFORMATION:
; APPLICANT: Bird, David MCK.
; APPLICANT: Wilson, Mark A.
; TITLE OF INVENTION: Nematode-Induced Genes in Tomato
; NUMBER OF SEQUENCES: 129
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Townsend and Townsend and Crew LLP
; STREET: Two Embarcadero Center, Eighth Floor
; CITY: San Francisco
; STATE: California
; COUNTRY: USA
; ZIP: 94111-3834
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent in Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/756,849
; FILING DATE: 26-NOV-1996
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/248,474
; FILING DATE: 25-MAY-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: Bastian, Kevin L.
; REGISTRATION NUMBER: 34,774
; REFERENCE/DOCKET NUMBER: 023070-053510US
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (415) 576-0200
; TELEFAX: (415) 576-0300
; INFORMATION FOR SEQ ID NO: 23:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 243 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; ORIGINAL SOURCE:
; ORGANISM: Lycopersicon esculentum cv 'Rutgers Large Red'
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: 1..243
; OTHER INFORMATION: /standard_name="DB# 139"
; US-08-756-849-23

Query Match          65.0%; Score 13; DB 3; Length 243;
Best Local Similarity 100.0%; Pred. No. 76;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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QY      3 TTAGAGGAGAACAA 15
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Db      133 TTAGAGGAGAACAA 121
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RESULT 28
US-09-702-705-1692/c
; Sequence 1692, Application US/09702705
; Patent No. 6504010
; GENERAL INFORMATION:
; APPLICANT: Wang, Tongtong
; APPLICANT: Bangur, Chaitanya S.
; APPLICANT: Lodes, Michael A.
; APPLICANT: Fanger, Gary
; APPLICANT: Vedavick, Tom
; APPLICANT: Carter, Derrick
; APPLICANT: Retter, Marc
; APPLICANT: Mannion, Jane
; APPLICANT: Fan, Liqun
```

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; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THE THERAPY AND
; FILE OF INVENTION: DIAGNOSIS OF LUNG CANCER
; FILE REFERENCE: 210121.478C14
; CURRENT APPLICATION NUMBER: US/09/702,705
; CURRENT FILING DATE: 2000-10-30
; NUMBER OF SEQ ID NOS: 1833
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 1692
; LENGTH: 450
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(450)
; OTHER INFORMATION: n = A,T,C or G
; US-09-702-705-1692

Query Match          65.0%; Score 13; DB 4; Length 450;
Best Local Similarity 100.0%; Pred. No. 73;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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QY      2 CTTAGAGGAGAACAA 14
      ||| ||| ||| ||| |||
Db      235 CTTAGAGGAGAACAA 223
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RESULT 29
US-09-736-457-1692/c
; Sequence 1692, Application US/09736457
; Patent No. 6509448
; GENERAL INFORMATION:
; APPLICANT: Wang, Tongtong
; APPLICANT: Bangur, Chaitanya S.
; APPLICANT: Lodes, Michael A.
; APPLICANT: Fanger, Gary
; APPLICANT: Vedavick, Tom
; APPLICANT: Carter, Derrick
; APPLICANT: Retter, Marc
; APPLICANT: Mannion, Jane
; APPLICANT: Fan, Liqun
; APPLICANT: Wang, Aljun
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THE THERAPY AND
; FILE REFERENCE: 210121.478C15
; CURRENT APPLICATION NUMBER: US/09/736,457
; CURRENT FILING DATE: 2000-12-13
; NUMBER OF SEQ ID NOS: 1864
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 1692
; LENGTH: 450
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(450)
; OTHER INFORMATION: n = A,T,C or G
; US-09-736-457-1692

Query Match          65.0%; Score 13; DB 4; Length 450;
Best Local Similarity 100.0%; Pred. No. 73;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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QY      2 CTTAGAGGAGAACAA 14
      ||| ||| ||| ||| |||
Db      235 CTTAGAGGAGAACAA 223
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```
RESULT 30
US-08-244-205-4/c
; Sequence 4, Application US/08244205
; Patent No. 5952544
; GENERAL INFORMATION:
; APPLICANT: Browse, John, Kinney, Anthony J.,
```

```

; APPLICANT: Pierce, John, Wierzbicki, Anna M.,
; APPLICANT: Yadav, Narendra S., Perez-Grau, Luis
; TITLE OF INVENTION: Fatty Acid Desaturase Genes
; TITLE OF INVENTION: from Plants
; NUMBER OF SEQUENCES: 32
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: E. I. du Pont de Nemours and Company
; STREET: 1007 Market Street
; CITY: Wilmington
; STATE: Delaware
; COUNTRY: U.S.A.
; ZIP: 19898
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: Macintosh
; OPERATING SYSTEM: Macintosh System, 6.0
; SOFTWARE: Microsoft Word, 4.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/244,205
; FILING DATE:
; CLASSIFICATION: 800
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 07/804,259
; FILING DATE: 4 DECEMBER 1991
; ATTORNEY/AGENT INFORMATION:
; NAME: Floyd, Linda A.
; REGISTRATION NUMBER: 33,692
; REFERENCE/DOCKET NUMBER: BB-1036-A
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (302) 992-4929
; TELEFAX: (302) 892-7949
; TELEX: 835420
; INFORMATION FOR SEQ ID NO: 4:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 1525 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: cDNA
; HYPOTHETICAL: NO
; ORIGINAL SOURCE:
; ORGANISM: Arabidopsis thaliana
; IMMEDIATE SOURCE:
; CLONE: PACF2-2
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 10..1350
; US-08-244-205-4

Query Match          65.0%; Score 13; DB 2; Length 1525;
Best Local Similarity 100.0%; Pred. No. 69;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY      6 GGAGGACAACTC 18
      |||||
      814 GGAGGACAACTC 802

Db

RESULT 31
PCT-US92-10284-4/c
; Sequence 4, Application PC/TUS9210284
; GENERAL INFORMATION:
; APPLICANT: Browe, John, Kinney, Anthony J.,
; APPLICANT: Pierce, John, Wierzbicki, Anna M.,
; APPLICANT: Yadav, Narendra S., Perez-Grau, Luis
; TITLE OF INVENTION: Fatty Acid Desaturase Genes
; TITLE OF INVENTION: from Plants
; NUMBER OF SEQUENCES: 32
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: E. I. du Pont de Nemours and Company
; STREET: 1007 Market Street
; CITY: Wilmington
; STATE: Delaware

```

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; COUNTRY: U.S.A.
; ZIP: 19898
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: Macintosh
; OPERATING SYSTEM: Macintosh System, 6.0
; SOFTWARE: Microsoft Word, 4.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: PCT/US92/10284
; FILING DATE: 19921203
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 07/804,259
; FILING DATE: 4 DECEMBER 1991
; ATTORNEY/AGENT INFORMATION:
; NAME: Floyd, Linda A.
; REGISTRATION NUMBER: 33,692
; REFERENCE/DOCKET NUMBER: BB-1036-A
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (302) 992-4929
; TELEFAX: (302) 892-7949
; TELEX: 835420
; INFORMATION FOR SEQ ID NO: 4:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 1525 base pairs
; TYPE: NUCLEIC ACID
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: cDNA
; HYPOTHETICAL: NO
; ORIGINAL SOURCE:
; ORGANISM: Arabidopsis thaliana
; IMMEDIATE SOURCE:
; CLONE: PACF2-2
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 10..1350
; PCT-US92-10284-4

Query Match          65.0%; Score 13; DB 5; Length 1525;
Best Local Similarity 100.0%; Pred. No. 69;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY      6 GGAGGACAACTC 18
      |||||
      814 GGAGGACAACTC 802

Db

RESULT 32
PCT-US94-01321-9/c
; Sequence 9, Application PC/TUS9401321
; GENERAL INFORMATION:
; APPLICANT:
; TITLE OF INVENTION: Altered linolenic and linoleic Acid Content
; TITLE OF INVENTION: in Plants
; NUMBER OF SEQUENCES: 72
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25 (EPO)
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: PCT/US94/01321
; FILING DATE: 04-FEB-1994
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/156551
; FILING DATE: 22-NOV-1993
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/014431
; FILING DATE: 05-FEB-1993
; INFORMATION FOR SEQ ID NO: 9:
; SEQUENCE CHARACTERISTICS:

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LENGTH: 1645 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: double  
TOPOLOGY: linear  
MOLECULE TYPE: CDNA  
FEATURE:  
NAME/KEY: CDS  
LOCATION: 125..1465  
PCT-US94-01321-9

Query Match 65.0%; Score 13; DB 5; Length 1645;  
Best Local Similarity 100.0%; Pred. No. 68;  
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 6 GGAGGAGCAAGTC 18  
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Db 929 GGAGGAGCAAGTC 917

RESULT 33  
US-09-389-956-79/c  
Sequence 79, Application US/09389956  
Patent No. 6586579  
GENERAL INFORMATION:  
APPLICANT: Huang, Shi  
TITLE OF INVENTION: PR-Domain Containing Nucleic Acids, Polypeptides,  
FILE REFERENCE: P-LJ 3611  
CURRENT APPLICATION NUMBER: US/09/389,956  
CURRENT FILING DATE: 1999-09-03  
NUMBER OF SEQ ID NOS: 93  
SOFTWARE: PatentIn Ver. 2.0  
SEQ ID NO 79  
LENGTH: 1970  
TYPE: DNA  
ORGANISM: Homo sapiens  
FEATURE:  
NAME/KEY: CDS  
LOCATION: (1)..(1596)  
US-09-389-956-79

Query Match 65.0%; Score 13; DB 4; Length 1970;  
Best Local Similarity 100.0%; Pred. No. 68;  
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 5 AGGAGGACAACT 17  
|||||  
Db 139 AGGAGGACAACT 127

RESULT 34  
US-10-020-079-39/c  
Sequence 39, Application US/10020079  
Patent No. 6579710  
GENERAL INFORMATION:  
APPLICANT: Turner, C. Alexander Jr.  
APPLICANT: Mathur, Brian  
TITLE OF INVENTION: No. 6579710e1 Human Kinases and Polynucleotides Encoding the Same  
FILE REFERENCE: LEX-0281-USA  
CURRENT APPLICATION NUMBER: US/10/020,079  
CURRENT FILING DATE: 2001-12-12  
PRIOR APPLICATION NUMBER: US 60/255,103  
PRIOR FILING DATE: 2000-12-12  
PRIOR APPLICATION NUMBER: US 60/289,422  
PRIOR FILING DATE: 2001-05-08  
NUMBER OF SEQ ID NOS: 40  
SOFTWARE: FastSeq for Windows Version 4.0  
SEQ ID NO 39  
LENGTH: 2517  
TYPE: DNA  
ORGANISM: homo sapiens  
US-10-020-079-39

Query Match 65.0%; Score 13; DB 4; Length 2517;  
Best Local Similarity 100.0%; Pred. No. 67;  
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 2 CTTAGGAGGAACA 14  
|||||  
Db 1296 CTTAGGAGGAACA 1284

RESULT 35  
US-10-020-079-37/c  
Sequence 37, Application US/10020079  
Patent No. 6579710  
GENERAL INFORMATION:  
APPLICANT: Turner, C. Alexander Jr.  
APPLICANT: Mathur, Brian  
TITLE OF INVENTION: No. 6579710e1 Human Kinases and Polynucleotides Encoding the Same  
FILE REFERENCE: LEX-0281-USA  
CURRENT APPLICATION NUMBER: US/10/020,079  
CURRENT FILING DATE: 2001-12-12  
PRIOR APPLICATION NUMBER: US 60/255,103  
PRIOR FILING DATE: 2000-12-12  
PRIOR APPLICATION NUMBER: US 60/289,422  
PRIOR FILING DATE: 2001-05-08  
NUMBER OF SEQ ID NOS: 40  
SOFTWARE: FastSeq for Windows Version 4.0  
SEQ ID NO 37  
LENGTH: 2556  
TYPE: DNA  
ORGANISM: homo sapiens  
US-10-020-079-37

Query Match 65.0%; Score 13; DB 4; Length 2556;  
Best Local Similarity 100.0%; Pred. No. 66;  
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 2 CTTAGGAGGAACA 14  
|||||  
Db 1335 CTTAGGAGGAACA 1323

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GenCore version 5.1.6  
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OM nucleic - nucleic search, using sw model

Run on: August 14, 2003, 20:57:44 ; Search time 126 Seconds  
(without alignments)  
428.482 Million cell updates/sec

Title: US-10-074-620-2  
Perfect score: 20  
Sequence: 1 cctagagagacagtcgcc 20

Scoring table: Oligo\_NUC  
Gapop 60.0 , Gapext 60.0

Searched: 2552756 seqs, 1349719017 residues

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Total number of hits satisfying chosen parameters: 5105512

Minimum DB seq length: 0

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Post-processing: Listing first 120 summaries

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25: /SIDSI1/gcgdata/geneseq/geneseq-emb1/NA2004.DAT:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

# SUMMARIES

Result No.	Score	* Query	Match Length	DB ID	Description
1	20	100.0	20	AAQ1012	Primer binding to
2	20	100.0	20	AAQ1012	EBNA 2 primer, Pos
3	20	100.0	20	AAQ1012	DNA sequence encod
4	20	100.0	20	AAQ1012	EBV gp350/220 CDNA
5	20	100.0	20	AAQ1012	Human pancreatic c
6	16	80.0	535	ABV96868	Human pancreatic c
7	16	80.0	556	ABV99010	Human pancreatic c
8	16	80.0	2376	AAH62708	Shrimp white spot
9	16	80.0	6572	AA158931	Human polynucleoti

9	16	80.0	7445	22	ABA09554	Human guanine nucl
10	16	80.0	7445	22	AAK51763	Human polynucleoti
11	16	80.0	7445	22	AAK52747	Human polynucleoti
12	16	80.0	7445	22	AA160717	Human polynucleoti
13	16	80.0	7843	23	AA582621	DNA encoding novel
14	16	80.0	9501	23	AA580083	Breast cancer asso
15	16	80.0	9783	24	ABK94929	Human novel polynu
16	16	80.0	9785	23	ABV22178	Human prostate exp
17	16	80.0	9785	23	ABV28017	Human prostate exp
18	16	80.0	180557	23	ABN85750	Human BAC clone Rp
19	16	80.0	305107	22	AAH62689	Shrimp white spot
20	15	75.0	123	21	AAQ20071	Human secreted pro
21	15	75.0	762	24	ABK99991	DNA encoding human
22	15	75.0	1691	24	ABZ66827	Arabidopsis thalia
23	15	75.0	17255	23	ABL29548	Drosophila melanog
24	15	75.0	53226	25	ABQ76886	Human G-protein co
25	14	70.0	125	15	AAQ76831	Human genome fragm
26	14	70.0	273	15	ABX30187	Human GDP-mannose
27	14	70.0	300	20	AAK98339	Human cancer cell
28	14	70.0	344	23	ABV06501	Human prostate exp
29	14	70.0	357	21	AAQ03189	Human secreted pro
30	14	70.0	408	22	AA581336	Novel human diagno
31	14	70.0	409	20	AAK55494	Rice Cmt1 homology
32	14	70.0	442	22	AA526323	Human CDNA encodin
33	14	70.0	442	25	ABX73664	Human novel polynu
34	14	70.0	467	23	ABV36454	Human prostate exp
35	14	70.0	477	22	ABA43976	Human breast cell
36	14	70.0	477	22	AAK02723	Human brain expres
37	14	70.0	477	22	AAK28164	Human bone marrow
38	14	70.0	477	24	ABSO2670	Human genome-deriv
39	14	70.0	484	24	ABN95344	Gene #1842 used to
40	14	70.0	484	24	ABN66312	Lung cancer relate
41	14	70.0	484	21	AAK79263	Human prostate exp
42	14	70.0	505	22	AAK35496	Human immune/haema
43	14	70.0	688	23	ABV24146	Human prostate exp
44	14	70.0	784	22	AA195704	Human neuroblastom
45	14	70.0	788	22	AA525868	Human CDNA encodin
46	14	70.0	788	25	ABX73209	Human novel polynu
47	14	70.0	788	25	AAH04091	Human CDNA clone (
48	14	70.0	854	21	AAH06579	Human immunogenic
49	14	70.0	854	22	AA563788	Human prostate exp
50	14	70.0	854	22	AAH93695	Human prostate-spe
51	14	70.0	854	22	AAH85009	Human prostate-spe
52	14	70.0	854	22	AAH02760	Prostate tumour an
53	14	70.0	854	22	AAH86992	Human P776P invent
54	14	70.0	854	22	ABJ95159	Human P776P CDNA s
55	14	70.0	854	25	ACA59536	Prostate cancer th
56	14	70.0	1144	23	ABL27133	Drosophila melanog
57	14	70.0	1245	18	AAH78772	DNA encoding Aquif
58	14	70.0	1245	25	ABX77244	Aquifex aspartate
59	14	70.0	1541	25	ABX70949	Novel human CDNA s
60	14	70.0	1632	22	AAH13876	Human CDNA sequenc
61	14	70.0	1701	21	AAQ44045	Zea mays DNA fragm
62	14	70.0	1763	22	AAH45659	Human protease reg
63	14	70.0	1821	21	AAQ51101	Arabidopsis thalia
64	14	70.0	3134	21	ABL27132	Drosophila melanog
65	14	70.0	3522	23	ABX71253	Human brain-deriv
66	14	70.0	3529	22	AAH76684	Human transcriptio
67	14	70.0	4619	23	ABX71386	Human testes-deriv
68	14	70.0	4809	22	AA563924	Human prostate exp
69	14	70.0	4809	22	AAH93831	Human prostate-spe
70	14	70.0	4809	24	ABL95295	Human prostate-spe
71	14	70.0	4809	25	ACA59772	Human P776P CDNA s
72	14	70.0	6250	22	AAJ36600	Human prostate cancer
73	14	70.0	6250	25	ABX59588	Human musculoskele
74	14	70.0	6251	22	AAJ36603	CDNA encoding nove
75	14	70.0	6251	25	ABX59591	Human musclelokele
76	14	70.0	9636	22	AAK90449	Human digestive sy
77	14	70.0	11446	22	AAK90444	Human digestive sy
78	14	70.0	16607	22	ABH18349	Human nervous syst
79	14	70.0	16607	22	AAH04678	Human reproductive
80	14	70.0	16607	23	ABL97585	Human testicular a
81	14	70.0	16738	22	AAK70864	Human immune/haema

C 82	14	70.0	24025	17	AAT17455	Mutated BRCA1 geno
C 83	14	70.0	24025	17	AAT17515	Mutated BRCA1 geno
C 84	14	70.0	24026	17	AAT17512	Mutated BRCA1 geno
C 85	14	70.0	24026	17	AAT17512	Mutated BRCA1 geno
C 86	14	70.0	24026	17	AAT17513	Mutated BRCA1 geno
C 87	14	70.0	24026	17	AAT17514	Mutated BRCA1 geno
C 88	14	70.0	24026	17	AAT17516	Mutated BRCA1 geno
C 89	14	70.0	24026	17	AAT17517	Mutated BRCA1 geno
C 90	14	70.0	24026	17	AAT17518	Mutated BRCA1 geno
C 91	14	70.0	24026	17	AAT17519	Mutated BRCA1 geno
C 92	14	70.0	24026	17	AAT17521	Mutated BRCA1 geno
C 93	14	70.0	24026	17	AAT17522	Mutated BRCA1 geno
C 94	14	70.0	24026	17	AAT17523	Mutated BRCA1 geno
C 95	14	70.0	24026	17	AAT17524	Mutated BRCA1 geno
C 96	14	70.0	24026	17	AAT17526	Mutated BRCA1 geno
C 97	14	70.0	24026	17	AAT17527	Mutated BRCA1 geno
C 98	14	70.0	24026	17	AAT17528	Mutated BRCA1 geno
C 99	14	70.0	24026	17	AAT17529	Mutated BRCA1 geno
C 100	14	70.0	24026	17	AAT17530	Mutated BRCA1 geno
C 101	14	70.0	24026	17	AAT17532	Mutated BRCA1 geno
C 102	14	70.0	24029	17	AAT17520	Mutated BRCA1 geno
C 103	14	70.0	24031	17	AAT17525	Mutated BRCA1 geno
C 104	14	70.0	25574	22	AAK05619	Human reproductive
C 105	14	70.0	25574	22	AAK79671	Human immune/haema
C 106	14	70.0	25574	22	AAK83760	Human immune/haema
C 107	14	70.0	25576	22	AAK05618	Human reproductive
C 108	14	70.0	25576	22	AAK79669	Human immune/haema
C 109	14	70.0	25576	22	AAK83758	Human immune/haema
C 110	14	70.0	25576	22	AAK83505	Human immune/haema
C 111	14	70.0	77425	24	ABK83502	Human cDNA differe
C 112	14	70.0	567571	25	AAD53224	Human chromosome 3
C 113	14	70.0	1691080	24	ABX08336	Human phosphodiester
C 114	13	65.0	114	19	AAV44485	tRNA-Lys3 pseudoge
C 115	13	65.0	126	18	AAT68500	Clone (individual)
C 116	13	65.0	126	18	AAT68881	Clone (individual)
C 117	13	65.0	242	22	ABAI1686	Human nervous syst
C 118	13	65.0	243	17	AAT09746	Tomato genomic DNA
C 119	13	65.0	286	24	ABL73501	Corn tassell-derive
C 120	13	65.0	294	24	ABL75621	Corn tassell-derive

## ALIGNMENTS

## RESULT 1

AAQ91012 standard; DNA, 20 BP.

AAQ91012;

01-FEB-1996 (first entry)

Primer binding to 3' end of EBV nuc antigen gene.

XX Primer; PCR; amplification; DNA polymerase; exonuclease; pfu; Tag;  
 KW Klenow fragment; T4; T7; Deep Vent; synthesis; mismatch; human; antibody;  
 heavy chain variable region; ss.

Synthetic.

MO9516028-A1.

15-JUN-1995.

07-DEC-1994; 94WO-US14065.

16-FEB-1994; 94US-0197791.

08-DEC-1993; 93US-0164290.

(STRA-) STRATAGENE.

Millimax RL, Sarge JA;

XX

WPI; 1995-224316/29.

Composn. useful for polynucleotide synthesis and cyclical

amplification - comprising a mixt. contg. an enzyme with 3'-5'

exo-nuclease activity and a DNA polymerase with less 3'-5'

Examples; Page 35; 66pp; English.

XX primers AAQ90984-Q91028 are examples of primers for testing a novel  
 CC composition for polynucleotide synthesis comprising a DNA polymerase  
 CC with high 3'-5' exonuclease activity in conjunction with a DNA polymerase  
 CC with less 3'-5' exonuclease activity, pref. Pfu and Tag DNA polymerases  
 CC respectively. Other DNA polymerases containing high 3'-5' exonuclease  
 CC activity include E. coli DNA polymerase I, Klenow fragment, T4, T7, Vent  
 CC or Deep Vent DNA polymerases. The use of a DNA polymerase with high  
 CC 3'-5' exonuclease activity is designed to overcome the inability of DNA  
 CC polymerases with low 3'-5' exonuclease activities to initiate synthesis  
 CC from primers containing 3' terminal mismatches, e.g. due to errors  
 CC introduced during a PCR cycle.  
 CC This primer binds to a region at the 5' end of the Epstein-Barr virus  
 CC nuc antigen gene.

Sequence 20 BP; 6 A; 6 C; 5 G; 3 T; 0 other;

Query Match 100.0%; Score 20; DB 16; Length 20;

Best Local Similarity 100.0%; Pred. No. 0.078; Mismatches 0; Gaps 0;

Matches 20; Conservative 0; Indels 0;

1 CCTTAGAGAGCAACATGCC 20

1 CCTTAGAGAGCAACATGCC 20

## RESULT 2

ABA00269 standard; DNA, 20 BP.

ABA00269;

29-NOV-2002 (first entry)

EBNA 2 primer, Position 90249;90269.

Primer; amplify; PCR; probe; detection; Epstein-Barr virus; EBV; ss.

Epstein-Barr virus.

WO200264842-A2.

22-AUG-2002.

13-FEB-2002; 2002WO-US04339.

13-FEB-2001; 2001US-268439P.

(CHIL-) CHILDRENS HOSPITAL RES FOUND.

Witte DP, Groen PA;

WPI; 2002-667015/71.

XX New compositions comprising nucleic acid sequences which specifically  
 PT hybridizes to Epstein-Barr virus (EBV) nucleic acid, for detecting EBV  
 PT in clinical specimens to determine patients at high risk of to  
 PT developing EBV infections

Claim 1; Page 44; 59pp; English.

XX The sequences given in ABA00268-75 are primers and probes which were  
 CC used in the compositions of the invention for the detection of  
 CC Epstein-Barr virus (EBV). The compositions comprise at least one  
 CC purified and isolated oligonucleotide consisting of a nucleic acid

CC	sequence which complements and specifically hybridizes to EBV nucleic
CC	acid. The oligonucleotide sequences and compositions comprising them
CC	are useful for detecting EBV in clinical specimens to determine
CC	patients who are at high risk to develop serious and costly medical
CC	complications, and allow for better clinical management of these
CC	patients by earlier recognition of their infection status. The
CC	oligonucleotide sequences may also be used to amplify EBV DNA
CC	sequences. The use of the oligonucleotide sequences in the assay for
CC	detecting EBV has a broad dynamic range of detection from less than
CC	10-100000000 copies of EBV DNA, is less labour intensive requiring only
CC	one reaction tube for the EBV determination, highly sensitive, accurate
CC	and has a rapid turn around time with assays that are completed,
CC	including amplification, probe specific hybridization, and calculation
CC	of copy number in less than 1 hour. The method may be adapted to
CC	automated systems.
CC	
XX	
XX	Sequence 20 BP; 6 A; 6 C; 5 G; 3 T; 0 other;
XX	
XX	Query Match 100.0%; Score 20; DB 24; Length 20;
XX	Best Local Similarity 100.0%; Pred. No. 0.078;
XX	Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY	1 CCTTAGGAGAACAAAGTCCC 20
Db	1 CCTTAGGAGAACAAAGTCCC 20
XX	
XX	RESULT 3
XX	AAN50114
XX	ID AAN50114 standard; DNA; 2721 BP.
XX	AAN50114;
XX	
XX	25-MAR-2003 (updated)
DT	17-OCT-1991 (first entry)
XX	
DE	DNA sequence encoding Epstein-Barr virus (EBV) outer surface protein.
XX	
XX	Epstein-Barr virus; antigen; vaccine; ss.
KW	
OS	Epstein-Barr virus.
XX	
XX	Key Location/Qualifiers
FT	mat_peptide 1..2721
FT	/*tag= a
FT	/label= EBV surface protein antigen
XX	
XX	EP151079-A.
XX	
XX	07-AUG-1985.
PD	
XX	
PF	28-JAN-1985; 85EP-0400141.
XX	
XX	23-JUL-1984; 84US-0633558.
PR	30-JAN-1984; 84US-0575352.
XX	
XX	(UYCH-) UNIV CHICAGO.
PA	
XX	
PI	Kleff E, Tanner J, Hummel M, Belssel C;
XX	
XX	WPI; 1985-191978/32.
DR	P-PSDB; AAP50073.
XX	
XX	
PT	New fragment of Epstein-Barr Virus DNA - useful in vector to
PT	express polypeptide for use in pregn. of vaccine against the
PT	virus and for use in diagnosis.
XX	
XX	Claim 1; Page 21-23; 26pp; English.
XX	
CC	The sequence encodes an outer surface viral protein of EBV, used
CC	to generate antidiodes reacting with the surface proteins of
CC	EBV-infected cells, and in the preparation of a vaccine against EBV.
CC	(Updated on 25-MAR-2003 to correct PA field.)

XX	Sequence	2721 BP; 762 A; 876 C; 557 G; 526 T; 0 other;
SQ		
Query Match	100.0%;	Score 20; DB 6; Length 2721;
Best Local Similarity	100.0%;	Pred. No. 0.058;
Matches	20; Conservative	0; Mismatches 0; Indels 0; Gaps 0;
OY	1 CCTTAGGAGCAACAAGTCCC 20 	
Db	1886 CCTTAGGAGCAACAAGTCCC 1905	
RESULT 4		
AAT04821	AAT04821 standard; cDNA; 5931 BP.	
XX	AAT04821;	
XX	18-JAN-1996 (first entry)	
DE	EBV gp350/220 cDNA.	
KM	EBV; gp350; gp220; gp350/gp220; non-splicing variant; vaccine; ds.	
XX	Epetein-Barr virus.	
OS		
Key	Location/Qualifiers	
FT	CDS	1014..3737
FT	/tag= a	1014..1067
FT	sig_peptide	/tag= b
FT	mat_peptide	1068..3734
FT	/tag= c	2514..2515
FT	misc_feature	/tag= d
FT	/function= splice donor site	
FT	/note= "bases 2513-2517 (AAGT) are replaced by GTCA in the non-splicing variant"	
FT	misc_feature	3105..3106
FT	/tag= e	
FT	/function= splice acceptor site	
FT	/note= "bases 3104-3107 (AGGT) are replaced by TGGA in the non-splicing variant"	
FT	polyA_signal	3742..3747
FT	/tag= f	
XX		
PN	W09528488-A1.	
XX		
PD	26-OCT-1995.	
XX		
PF	13-APR-1995; 95WO-US04611.	
XX		
PR	18-APR-1994; 94US-0229291.	
XX	(AVIR-) AVIRON.	
PA		
XX		
PI	Jackman WT, Spaete R;	
XX		
DR	WPI, 1995-373802/48.	
XX	P-P8SDB; AAR80144.	
PT	New DNA encoding a homogeneous gp350 protein - can be used for preventing and treating Epstein-Barr virus-related diseases or conditions	
PS	Claim 2; Fig.1; 61pp; English.	
CC	The donor and acceptor splice sites of the EBV gene encoding gp350/220 are mutated by replacement of native nucleotides by non-native nucleotides, without altering the encoded amino acid sequence, resulting in elimination of gp220 prodn. Recombinant homogeneous gp350, useful in vaccines, is expressed in mammalian or insect cell hosts.	
CC		

XX SQ Sequence 5931 BP; 1453 A; 1782 C; 1437 G; 1259 T; 0 other;  
Query Match 100.0%; Score 20; DB 16; Length 5931;  
Best Local Similarity 100.0%; Pred. No. 0.055;  
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
Oy 1 CCTAGAGAGAACAGTCCC 20  
Db 2899 CTTAGAGAGAACAGTCCC 2918  
RESULT 5  
ABV96868  
ID ABV96868 standard; cDNA; 535 BP.  
XX AC ABV96868;  
XX DT 14-JAN-2003 (first entry)  
XX DE Human pancreatic cancer expressed cDNA SEQ ID NO 2276.  
XX KM Human; pancreas; cancer; gene therapy; vaccine; immunostimulant;  
XX KW cytosolic; tumour; gene; ss.  
XX OS Homo sapiens.  
XX PN WO200260317-A2.  
XX PD 08-AUG-2002.  
XX PF 30-JAN-2002; 2002WO-US02781.  
XX PR 30-JAN-2001; 2001US-265305P.  
XX PR 31-JAN-2001; 2001US-265682P.  
XX PR 09-FEB-2001; 2001US-267568P.  
XX PR 21-MAR-2001; 2001US-278651P.  
XX PR 28-APR-2001; 2001US-287112P.  
XX PR 16-MAY-2001; 2001US-291631P.  
XX PR 12-JUL-2001; 2001US-305484P.  
XX PR 20-AUG-2001; 2001US-313999P.  
XX PR 27-NOV-2001; 2001US-333626P.  
XX PA (CORI-) CORIXA CORP.  
XX PI Benson DR, Kalos MD, Lodes MJ, Persing DH, Hepler WT, Jiang Y;  
XX DR WPI; 2002-627435/67.  
XX PT New isolated polynucleotide and pancreatic tumor polypeptides, useful  
XX PT for diagnosing, preventing and/or treating cancer, particularly  
XX PT pancreatic cancer -  
XX PS Claim 1; SEQ ID NO 2276; 300pp + Sequence Listing; English.  
XX XX  
XX The invention relates to an isolated polynucleotide (I) comprising: (a)  
XX any of a group of over 4000 nucleotide sequences (ABV94628-ABV9145);  
XX (b) complements of (a); (c) sequences consisting of at least 20  
XX contiguous residues of (a); (d) sequences that hybridize to (a), under  
XX moderately stringent conditions; (e) sequences having at least 75% or 90%  
XX identity to (a); or (f) degenerate variants of (a). Polypeptides  
XX (ABP68596-ABP68637) encoded by (I) and oligonucleotide can be used to  
XX detect cancer in a patient and compositions comprising polypeptides,  
XX polynucleotides, antibodies, fusion proteins, T cell populations and  
XX antigen presenting cells expressing the polypeptide are useful in  
XX treating pancreatic cancer and stimulating an immune response. The  
XX polynucleotides can be used as probes or primers for nucleic acid  
XX hybridisation, in the design and preparation of ribozyme molecules for  
XX inhibiting expression of the tumour polypeptides and proteins in the  
XX tumour cells, in vaccines and for gene therapy.  
XX Note: The sequence data for this patent did not form part of the printed  
XX specification, but was obtained in electronic format directly from WIPO  
XX at ftp.wipo.int/pub/published\_pct\_sequences.

XX SQ Sequence 535 BP; 171 A; 112 C; 104 G; 148 T; 0 other;  
Query Match 80.0%; Score 16; DB 24; Length 535;  
Best Local Similarity 100.0%; Pred. No. 10;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
Oy 2 CTTAGAGAGAACAGT 17  
Db 87 CTTAGAGAGAACAGT 102  
RESULT 6  
ABV99010  
ID ABV99010 standard; cDNA; 556 BP.  
XX AC ABV99010;  
XX DT 14-JAN-2003 (first entry)  
XX DE Human pancreatic cancer expressed cDNA SEQ ID NO 4418.  
XX KM Human; pancreas; cancer; gene therapy; vaccine; immunostimulant;  
XX KW cytosolic; tumour; gene; ss.  
XX OS Homo sapiens.  
XX PN WO200260317-A2.  
XX PD 08-AUG-2002.  
XX PF 30-JAN-2002; 2002WO-US02781.  
XX PR 30-JAN-2001; 2001US-265305P.  
XX PR 31-JAN-2001; 2001US-265682P.  
XX PR 09-FEB-2001; 2001US-267568P.  
XX PR 21-MAR-2001; 2001US-278651P.  
XX PR 28-APR-2001; 2001US-287112P.  
XX PR 16-MAY-2001; 2001US-291631P.  
XX PR 12-JUL-2001; 2001US-305484P.  
XX PR 20-AUG-2001; 2001US-313999P.  
XX PR 27-NOV-2001; 2001US-333626P.  
XX PA (CORI-) CORIXA CORP.  
XX PI Benson DR, Kalos MD, Lodes MJ, Persing DH, Hepler WT, Jiang Y;  
XX DR WPI; 2002-627435/67.  
XX PT New isolated polynucleotide and pancreatic tumor polypeptides, useful  
XX PT for diagnosing, preventing and/or treating cancer, particularly  
XX PT pancreatic cancer -  
XX PS Claim 1; SEQ ID NO 4418; 300pp + Sequence Listing; English.  
XX XX  
XX The invention relates to an isolated polynucleotide (I) comprising: (a)  
XX any of a group of over 4000 nucleotide sequences (ABV94628-ABV9145);  
XX (b) complements of (a); (c) sequences consisting of at least 20  
XX contiguous residues of (a); (d) sequences that hybridize to (a), under  
XX moderately stringent conditions; (e) sequences having at least 75% or 90%  
XX identity to (a); or (f) degenerate variants of (a). Polypeptides  
XX (ABP68596-ABP68637) encoded by (I) and oligonucleotide can be used to  
XX detect cancer in a patient and compositions comprising polypeptides,  
XX polynucleotides, antibodies, fusion proteins, T cell populations and  
XX antigen presenting cells expressing the polypeptide are useful in  
XX treating pancreatic cancer and stimulating an immune response. The  
XX polynucleotides can be used as probes or primers for nucleic acid  
XX hybridisation, in the design and preparation of ribozyme molecules for  
XX inhibiting expression of the tumour polypeptides and proteins in the  
XX tumour cells, in vaccines and for gene therapy.  
XX Note: The sequence data for this patent did not form part of the printed  
XX specification, but was obtained in electronic format directly from WIPO  
XX at ftp.wipo.int/pub/published\_pct\_sequences.

XX Sequence 556 BP; 173 A; 117 C; 114 G; 152 T; 0 other;  
SQ  
Query Match 80.0%; Score 16; DB 24; Length 556;  
Best Local Similarity 100.0%; Pred. No. 10;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
OY 2 CTTAGAGAGAACAGT 17  
DB 108 CTTAGAGAGAACAGT 123  
RESULT 7  
AAH62708  
ID AAH62708 standard; DNA; 2376 BP.  
XX  
AC AAH62708;  
XX  
XX 11-SEP-2001 (first entry)  
XX  
XX Shrimp white spot Bacilliform virus (WSBV) gene 19.  
XX  
XX Shrimp white spot Bacilliform virus; WSBV; diagnosis; viral infection;  
XX  
XX antiviral agent; gene expression; antisense construct;  
XX  
XX transgenic viral resistant shrimp; ds.  
XX  
XX White spot syndrome virus.  
XX  
XX WO200138351-A2.  
XX  
XX 31-MAY-2001.  
XX  
XX 08-NOV-2000; 2000WO-US28888.  
XX  
XX 24-NOV-1999; 99CN-0124717.  
XX  
XX (PENY-) PE CORP NY.  
XX  
XX (THIR-) THIRD INST OCEANOGRAPHY STATE OCEANI C A.  
XX  
XX (SINO-) SINOGENOMAX CO LTD.  
XX  
XX Xu X, Yang F, He J, Pham L, He M, Ye Y, Shen Y, Kodira C;  
XX  
XX WPI; 2001-355877/37.  
XX  
XX P-PSDB; AAG84928.  
XX  
XX Primary nucleotide sequence of the shrimp white spot Bacilliform virus  
XX  
XX (WSBV), useful for producing viral polypeptides that can be used to  
XX  
XX screen for agents that are useful for treating WSBV infection -  
XX  
XX  
XX Claim 4; Figure 3; 626bp; English.  
XX  
XX The invention provides the primary nucleotide sequence of the WSBV genome  
XX  
XX (AAH62689), predicted transcript sequences (AAH62689-AAH62839) and  
XX  
XX encoded proteins (AAG84910-AAG85051) and oligonucleotide sequences  
XX  
XX (AAH6840-63160) suitable for use as primers or probes. The nucleic acid  
XX  
XX molecules and proteins of the invention are useful for diagnosis and  
XX  
XX monitoring viral infection. In screens for antiviral agents and for  
XX  
XX monitoring viral gene expression or activity during a treatment regimen.  
XX  
XX The nucleic acid molecules are also useful as antisense constructs to  
XX  
XX control viral gene expression in infected cells and tissues and to create  
XX  
XX transgenic viral resistant shrimp.  
XX  
XX Sequence 2376 BP; 787 A; 467 C; 526 G; 596 T; 0 other;  
SQ  
Query Match 80.0%; Score 16; DB 22; Length 2376;  
Best Local Similarity 100.0%; Pred. No. 9.3;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

RESULT 8  
AA158931  
ID AA158931 standard; cDNA; 6572 BP.  
XX  
XX AA158931;  
XX  
XX 22-OCT-2001 (first entry)  
XX  
XX  
XX Human polynucleotide SEQ ID NO 1134.  
XX  
XX  
XX Human; nootropic; immunosuppressant; cytostatic; gene therapy; cancer;  
XX  
XX peripheral nervous system; neuropathy; central nervous system; CNS;  
XX  
XX Alzheimer's; Parkinson's disease; Huntington's disease; haemostatic;  
XX  
XX amyotrophic lateral sclerosis; Shy-Drager Syndrome; chemotactic;  
XX  
XX chemokine; thrombolytic; drug screening; arthritis; inflammation;  
XX  
XX leukaemia; ss.  
XX  
XX Homo sapiens.  
XX  
XX WO200153312-A1.  
XX  
XX 26-JUL-2001.  
XX  
XX 26-DEC-2000; 2000WO-US34263.  
XX  
XX 21-JAN-2000; 2000US-0488725.  
XX  
XX 25-APR-2000; 2000US-0552317.  
XX  
XX 09-JUL-2000; 2000US-0598042.  
XX  
XX 19-JUL-2000; 2000US-0620312.  
XX  
XX 03-AUG-2000; 2000US-0653450.  
XX  
XX 14-SEP-2000; 2000US-0662191.  
XX  
XX 19-OCT-2000; 2000US-0693036.  
XX  
XX 29-NOV-2000; 2000US-0727344.  
XX  
XX (HYSE-) HYSEQ INC.  
XX  
XX Tang YT, Liu C, Asundi V, Chen R, Ma Y, Qian XB, Ren F, Wang D;  
XX  
XX Wang Z, Wehrman T, Xu C, Xue AJ, Yang Y, Zhang J;  
XX  
XX Zhao QA, Zhou P, Goodrich R, Drmanac RT;  
XX  
XX WPI; 2001-442253/47.  
XX  
XX P-PSDB; AAM39775.  
XX  
XX Novel nucleic acids and polypeptides, useful for treating disorders  
XX  
XX such as central nervous system injuries -  
XX  
XX  
XX Claim 1; SEQ ID NO 1134; 10078bp; English.  
XX  
XX The invention relates to human nucleic acids (AA157798-AA161369) and  
XX  
XX the encoded polypeptides (AAM38642-AAM42213) with nootropic,  
XX  
XX immunosuppressant and cytostatic activity. The polynucleotides are useful  
XX  
XX in gene therapy. A composition containing a polypeptide or polynucleotide  
XX  
XX of the invention may be used to treat diseases of the peripheral nervous  
XX  
XX system, such as peripheral nervous injuries, peripheral neuropathy and  
XX  
XX localized neuropathies and central nervous system diseases, such as  
XX  
XX Alzheimer's, Parkinson's disease, Huntington's disease, amyotrophic  
XX  
XX lateral sclerosis, and Shy-Drager Syndrome. Other uses include the  
XX  
XX utilisation of the activities such as: Immune system suppression,  
XX  
XX Activin/inhibin activity, chemotactic/chemokinetic activity, haemostatic  
XX  
XX and thrombolytic activity, cancer diagnosis and therapy, drug screening,  
XX  
XX assays for receptor activity, arthritis and inflammation, leukaemias and  
XX  
XX C.N.S disorders.  
XX  
XX Note: The sequence data for this patent did not form part of the printed  
XX  
XX specification.  
SQ  
Sequence 6572 BP; 1929 A; 1256 C; 1418 G; 1969 T; 0 other;  
Query Match 80.0%; Score 16; DB 22; Length 6572;  
Best Local Similarity 100.0%; Pred. No. 8.8;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
OY 2 CTTAGAGAGAACAGT 17  
DB 108 CTTAGAGAGAACAGT 123

Db 4524 CTTAGGAGAACACT 4539

RESULT 9

ID ABA09554 standard; cDNA, 7445 BP.

XX ABA09554;

XX 11-JAN-2002 (first entry)

XX Human guanine nucleotide exchange factor homologue cDNA, SEQ ID NO:1330.

XX

XX Human: cytokine; cell proliferation; tissue growth; immunomodulatory; growth factor; haematopoiesis regulation; tissue growth; immunomodulatory; actinin; inhibin; chemotaxis; chemokinesis; thrombolysis; oncogenesis; myeloid cell disorder; lymphoid cell disorder; asthma; arthritis; chronic inflammatory condition; proliferative retinopathy; atherosclerosis; coronary heart disease; arterial ischaemia; bone disorder; osteoporosis; vascular growth disorder; tissue regeneration; wound healing; infection; immune disorder; cell culture; drug screening; gene therapy; antiinflammatory; antiaesthetic; antiarthritic; haemostatic; antiarteriosclerotic; cycstatic; osteopathic; vasotropic; cardiant; virucide; antibacterial; antifungal; vulnerary; antitumor; ss.

XX

XX Homo sapiens.

XX WO200157188-A2.

XX

XX 09-AUG-2001.

XX

XX 05-FEB-2001; 2001WO-US03800.

XX

XX 03-FEB-2000; 2000US-0496914.

XX 27-APR-2000; 2000US-0560875.

XX (HYSE-) HYSEQ INC.

XX Tang YT, Liu C, Drmanac RT;

XX WPI; 2001-457740/49.

XX P-PSDB; ABB12310.

XX Human proteins and DNA encoding sequences useful for preventing treating or ameliorating a medical condition in a mammalian subject e.g. arthritis and cancer -

XX

XX Claim 1; Page 992-994; 1963jp; English.

XX

XX Sequences ABB10991-ABB12330 represent 1350 novel human polypeptides, and sequences ABA08225-ABA09574 represent nucleic acids encoding them. The invention also relates to vectors and recombinant host cells comprising a nucleotide of the invention, methods of producing the novel polypeptides, antibodies against the polypeptides, methods of detecting the nucleotides or polypeptides in a sample, and methods of identifying compounds which bind to polypeptides of the invention. Although novel, many of the polypeptides of the invention have homology to known proteins, thereby giving an insight into their probable biological activities, and hence potential therapeutic applications. The polypeptides of the invention may have various activities, including cytokine, cell proliferation or cell differentiation activities; stem cell growth factor activity; haematopoiesis regulatory activity; tissue growth activity; immunomodulatory activity; activin- or inhibin-related activities; chemotactic or chemokinetic activities; haemostatic, thrombotic or thrombolytic activities; receptor or ligand activities; or may be involved in oncogenesis, cancer cell proliferation or metastasis. Depending on their biological activities, polypeptides and nucleotides of the invention are useful for preventing, treating or ameliorating medical conditions, e.g., by protein or gene therapy. Such conditions include cancers, haematopoietic disorders (e.g., myeloid or lymphoid cell disorders), chronic inflammatory conditions (e.g., asthma or arthritis),

CC proliferative retinopathy, atherosclerosis, coronary heart disease, CC arterial ischaemia, bone disorders (e.g., osteoporosis), and abnormal CC vascular growth. Polypeptides involved with tissue regeneration and CC repair (or nucleic acids encoding them) may be used to promote wound CC healing (e.g., of burns, incisions and ulcers), while those with CC immunomodulatory activities may be used in the treatment of viral, CC bacterial and fungal infections in addition to immune disorders. CC Polypeptides with growth factor activity may be used in cell cultures to CC promote cell growth. For example, such polypeptides may be used to CC manipulate stem cells in culture to give rise to neuroepithelial cells CC that can be used to augment or replace cells damaged by illness, CC autoimmune disease or accidental damage. The polypeptides and nucleotides CC may also be used in the diagnosis of the above conditions, and in drug CC screening techniques. The present sequence represents a cDNA encoding a CC novel human polypeptide of the invention.

XX

XX Sequence 7445 BP; 2177 A; 1404 C; 1582 G; 2282 T; 0 other;

XX

XX Query Match 80.0%; Score 16; DB 22; Length 7445;

XX Best Local Similarity 100.0%; Pred. No. 8.7;

XX Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

XX

XX 2 CTTAGGAGAACACT 17

XX

XX Db 4492 CTTAGGAGAACACT 4507

XX

XX RESULT 10

XX AAK51763

XX ID AAK51763 standard; cDNA, 7445 BP.

XX

XX AAK51763;

XX

XX 06-NOV-2001 (first entry)

XX

XX Human polynucleotide SEQ ID NO 308.

XX

XX Human: cytokine; cell proliferation; cell differentiation; gene therapy; tissue growth factor; stem cell growth factor; haematopoiesis; tissue growth factor; immunomodulatory; cancer; leukaemia; nervous system disorder; arthritis; inflammation; ss.

XX

XX Homo sapiens.

XX WO200157190-A2.

XX

XX 09-AUG-2001.

XX

XX 05-FEB-2001; 2001WO-US04098.

XX

XX 03-FEB-2000; 2000US-0496914.

XX 27-APR-2000; 2000US-0560875.

XX 20-JUN-2000; 2000US-0598075.

XX 19-JUN-2000; 2000US-0620325.

XX 01-SEP-2000; 2000US-0654936.

XX 15-SEP-2000; 2000US-0663561.

XX 20-OCT-2000; 2000US-0693325.

XX 30-NOV-2000; 2000US-0728422.

XX

XX (HYSE-) HYSEQ INC.

XX

XX Tang YT, Liu C, Drmanac RT, Asundi V, Zhou P, Xu C, Cao Y, Ma Y;

XX Zhao YA, Wang D, Wang J, Zhang J, Ren F, Chen R, Wang ZW;

XX Xue AJ, Yang Y, Wehrman T, Goodrich R;

XX WPI; 2001-476283/51.

XX P-PSDB; AAM78630.

XX Nucleic acids encoding polypeptides with cytokine-like activities, PT useful in diagnosis and gene therapy -

XX

XX Claim 1; Page 1279-1284; 6221jp; English.

CC The invention relates to polynucleotides (AAK51456-AAK53435) and the  
CC encoded polypeptides (AAM78323-AAM80302) that exhibit activity elating to  
CC cytokine, cell proliferation or cell differentiation or which may induce  
CC production of other cytokines in other cell populations. The  
CC polynucleotides and polypeptides are useful in gene therapy, vaccines or  
CC peptide therapy. The polypeptides have various cytokine-like activities,  
CC e.g. stem cell growth factor activity, haematopoietic regulating  
CC activity, tissue growth factor activity, immunomodulatory activity and  
CC activity/inhibin activity and may be useful in the diagnosis and/or  
CC treatment of cancer, leukaemia, nervous system disorders, arthritis and  
CC inflammation.  
CC Note: Records for SEQ ID NO 2110 (AAK52581), 2111 (AAK52582) and 3666  
CC (AAM80020) are omitted as the relevant pages from the sequence listing  
CC were missing at the time of publication.

SO Sequence 7445 BP; 2179 A; 1403 C; 1584 G; 2279 T; 0 other;

Query Match 80.0%; Score 16; DB 22; Length 7445;

Best Local Similarity 100.0%; Pred. No. 8.7;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 CTTAGGAGAACAACT 17  
DB 4481 CTTAGGAGAACAACT 4496

## RESULT 11

AAK52747  
ID AAK52747 standard; cDNA; 7445 BP.

AC AAK52747;

DT 06-NOV-2001 (first entry)

DE Human polynucleotide SEQ ID NO 2276.

XX Human; cytokine; cell proliferation; cell differentiation; gene therapy;

KM vaccine; peptide therapy; stem cell growth factor; haematopoiesis;

KM tissue growth factor; immunomodulatory; cancer; leukaemia;

KM nervous system disorder; arthritis; inflammation; ss.

XX Homo sapiens.

XX WO200157190-A2.

PD 09-AUG-2001.

PF 05-FEB-2001; 2001WO-US04098.

XX 03-FEB-2000; 2000US-0496914.

PR 27-APR-2000; 2000US-0560875.

PR 20-JUN-2000; 2000US-0598075.

PR 19-JUL-2000; 2000US-0620325.

PR 01-SEP-2000; 2000US-0654936.

PR 15-SEP-2000; 2000US-0663561.

PR 20-OCT-2000; 2000US-0693325.

PR 30-NOV-2000; 2000US-0728422.

XX (HYSE-) HYSEQ INC.

XX Tang YT, Liu C, Drmanac RT, Asundi V, Zhou P, Xu C, Cao Y, Ma Y,

PI Zhao QA, Wang D, Wang J, Zhang J, Ren F, Chen R, Wang ZW;

PI Xue AJ, Yang Y, Wejthman T, Goodrich R;

XX WPI; 2001-476283/51.

DR P-PSDB; AAM79614.

XX Nucleic acids encoding polypeptides with cytokine-like activities,

PT useful in diagnosis and gene therapy -

PS Claim 1; Page 4614-4616; 6221pp; English.

XX The invention relates to polynucleotides (AAK51456-AAK53435) and the

CC encoded polypeptides (AAM78323-AAM80302) that exhibit activity elating to  
CC cytokine, cell proliferation or cell differentiation or which may induce  
CC production of other cytokines in other cell populations. The  
CC polynucleotides and polypeptides are useful in gene therapy, vaccines or  
CC peptide therapy. The polypeptides have various cytokine-like activities,  
CC e.g. stem cell growth factor activity, haematopoietic regulating  
CC activity, tissue growth factor activity, immunomodulatory activity and  
CC activity/inhibin activity and may be useful in the diagnosis and/or  
CC treatment of cancer, leukaemia, nervous system disorders, arthritis and  
CC inflammation.  
CC Note: Records for SEQ ID NO 2110 (AAK52581), 2111 (AAK52582) and 3666  
CC (AAM80020) are omitted as the relevant pages from the sequence listing  
CC were missing at the time of publication.

SO Sequence 7445 BP; 2177 A; 1404 C; 1582 G; 2282 T; 0 other;

Query Match 80.0%; Score 16; DB 22; Length 7445;

Best Local Similarity 100.0%; Pred. No. 8.7;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 CTTAGGAGAACAACT 17  
DB 4492 CTTAGGAGAACAACT 4507

## RESULT 12

AAI60717  
ID AAI60717 standard; cDNA; 7445 BP.

AC AAI60717;

DT 22-OCT-2001 (first entry)

DE Human polynucleotide SEQ ID NO 4706.

XX Human; nootropic; immunosuppressant; cytostatic; gene therapy; cancer;

KM peripheral nervous system; neuropathy; central nervous system; CNS;

KM Alzheimer's; Parkinson's disease; Huntington's disease; haemostatic;

KM amyotrophic lateral sclerosis; Shy-Drager Syndrome; chemotactic;

KM chemokinetic; thrombolytic; drug screening; arthritis; inflammation;

KM leukaemia; ss.

XX Homo sapiens.

XX WO200153312-A1.

PD 26-JUL-2001.

PF 26-DEC-2000; 2000WO-US34263.

XX 21-JAN-2000; 2000US-0488725.

PR 25-APR-2000; 2000US-0552317.

PR 09-JUL-2000; 2000US-0598042.

PR 19-JUL-2000; 2000US-0620312.

PR 03-AUG-2000; 2000US-0653450.

PR 14-SEP-2000; 2000US-0662191.

PR 19-OCT-2000; 2000US-0693036.

PR 29-NOV-2000; 2000US-0727344.

XX (HYSE-) HYSEQ INC.

XX Tang YT, Liu C, Drmanac RT, Asundi V, Zhou P, Xu C, Cao Y, Ma Y,

PI Wang J, Wang Z, Wejthman T, Xu C, Xue AJ, Yang Y, Zhang J,

PI Zhao QA, Zhou P, Goodrich R, Drmanac RT;

XX WPI; 2001-442253/47.

DR P-PSDB; AAM41561.

XX Novel nucleic acids and polypeptides, useful for treating disorders

PT such as central nervous system injuries -

PS Claim 1; SEQ ID NO 4706; 10078pp; English.

XX

CC The invention relates to human nucleic acids (AA157798-AA161369) and  
CC the encoded polypeptides (AA038642-AA042213) with neurotropic.  
CC immunosuppressant and cytostatic activity. The polynucleotides are useful  
CC in gene therapy. A composition containing a polypeptide or polynucleotide  
CC of the invention may be used to treat diseases of the peripheral nervous  
CC system, such as peripheral nervous injuries, peripheral neuropathy and  
CC localised neuropathies and central nervous system diseases, such as  
CC Alzheimer's, Parkinson's disease, Huntington's disease, amyotrophic  
CC lateral sclerosis, and Shy-Drager Syndrome. Other uses include the  
CC utilisation of the activities such as: Immune system suppression,  
CC Activin/inhibin activity, chemotactic/chemokinetic activity, haemostatic  
CC and thrombolytic activity, cancer diagnosis and therapy, drug screening,  
CC assays for receptor activity, arthritis and inflammation, leukaemias and  
CC C.N.S disorders.  
CC Note: The sequence data for this patent did not form part of the printed  
CC specification.

XX  
XX  
SQ Sequence 7445 BP; 2177 A; 1404 C; 1582 G; 2282 T; 0 other;  
Query Match 80.0%; Score 16; DB 22; Length 7445;  
Best Local Similarity 100.0%; Pred. No. 8.7;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 CTTAGGAGGAAACAAGT 17  
|||||  
Db 4492 CTTAGGAGGAAACAAGT 4507

RESULT 13  
AAS82621  
ID AAS82621 standard; cDNA; 7843 BP.  
XX  
XX AC AAS82621;  
XX  
XX 13-FEB-2002 (first entry)  
XX  
XX DNA encoding novel human diagnostic protein #18425.  
XX  
XX Human: chromosome mapping; gene mapping; gene therapy; forensic;  
XX food supplement; medical imaging; diagnostic; genetic disorder; ss.  
XX  
XX Homo sapiens.  
XX OS  
XX WO200175067-A2.  
XX  
XX 11-OCT-2001.  
XX  
XX 30-MAR-2001; 2001WO-US08631.  
XX  
XX 31-MAR-2000; 2000US-0540217.  
XX PR 23-AUG-2000; 2000US-0649167.  
XX  
XX (HYSE-) HYSEQ INC.  
XX  
XX  
XX Dmanac RT, Liu C, Tang YT;  
XX  
XX WPI: 2001-639362/73.  
XX DR P-FSDB; ABG18434.  
XX  
XX New isolated polynucleotide and encoded polypeptides, useful in  
XX PT diagnostics, forensics, gene mapping, identification of mutations  
XX PT responsible for genetic disorders or other traits and to assess  
XX PT biodiversity -  
XX  
XX Claim 1; SEQ ID No 18425; 103bp; English.

CC The invention relates to isolated polynucleotide (I) and  
CC polypeptide (II) sequences. (I) is useful as hybridisation probes,  
CC polymerase chain reaction (PCR) primers, oligomers and for chromosome  
CC and gene mapping, and in recombinant production of (II). The  
CC polynucleotides are also used in diagnostics as expressed sequence tags  
CC for identifying expressed genes. (I) is useful in gene therapy techniques  
CC to restore normal activity of (II) or to treat disease states involving

CC (II). (II) is useful for generating antibodies against it, detecting or  
CC quantitating a polypeptide in tissue, as molecular weight markers and as  
CC a food supplement. (II) and its binding partners are useful in medical  
CC imaging of sites expressing (II). (I) and (II) are useful for treating  
CC disorders involving aberrant protein expression or biological activity.  
CC The polypeptide and polynucleotide sequences have applications in  
CC diagnostics, forensics, gene mapping, identification of mutations  
CC responsible for genetic disorders or other traits to assess biodiversity  
CC and to produce other types of data and products dependent on DNA and  
CC amino acid sequences. AAS64197-AAS94564 represent novel human  
CC diagnostic coding sequences of the invention.  
CC Note: The sequence data for this patent did not appear in the printed  
CC specification, but was obtained in electronic format directly from WIPO  
CC at ftp.wipo.int/pub/published\_pct\_sequences.

XX  
XX  
SQ Sequence 7843 BP; 2262 A; 1492 C; 1682 G; 2403 T; 4 other;  
Query Match 80.0%; Score 16; DB 23; Length 7843;  
Best Local Similarity 100.0%; Pred. No. 8.7;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 CTTAGGAGGAAACAAGT 17  
|||||  
Db 4761 CTTAGGAGGAAACAAGT 4776

RESULT 14  
ACC50083  
ID ACC50083 standard; cDNA; 9501 BP.  
XX  
XX AC ACC50083;  
XX  
XX 12-JUN-2003 (first entry)  
XX  
XX Breast cancer associated cDNA sequence SEQ ID NO:14.  
XX  
XX Human; breast cancer; cytostatic; gene therapy; gene; ss.  
XX  
XX Homo sapiens.  
XX OS  
XX WO2003004989-A2.  
XX  
XX 16-JAN-2003.  
XX  
XX 21-JUN-2002; 2002WO-US19669.  
XX  
XX 21-JUN-2001; 2001US-299887P.  
XX PR 27-JUN-2001; 2001US-301572P.  
XX PR 18-JUN-2001; 2001US-306501P.  
XX PR 25-SEP-2001; 2001US-325002P.  
XX PR 05-MAR-2002; 2002US-362585P.  
XX PR 14-MAY-2002; 2002US-380391P.  
XX  
XX (MTLL-) MILENITUM PHARM INC.  
XX  
XX  
XX Lillie J, Gannavarapu M, Glact K, Hoersh S, Kamatkar S, Mertens M;  
XX PI Monahan JB, Myer V, Wang Y, Xu Y, Zhao X, Meyers RJ, Bast RC;  
XX PI Hortobagyi GN, Puertal L, Meric F, Sahin A, Mills GB;  
XX  
XX WPI: 2003-210381/20.  
XX DR P-FSDB; ABR47392.  
XX  
XX Breast cancer diagnosis or treatment by comparing the level of  
XX PT expression of a marker in a patient sample with that in the control  
XX PT non-breast cancer sample -  
XX  
XX Claim 1; SEQ ID 14; 128bp; English.

CC The present invention describes a method for assessing whether a patient  
CC is afflicted with breast cancer. The method comprises comparing the level  
CC of expression of a marker (gene/polypeptide see ACC50076 to ACC50334 and  
CC ABR447386 to ABR47632) in a patient sample and the normal level of  
CC expression of the marker in a control non-breast cancer sample, where a



CC significant increase in the level of expression of the marker in the  
CC patient sample and the normal level is an indication that the patient is  
CC afflicted with breast cancer. The breast cancer associated sequences  
CC from the present invention have cytostatic activities and can be used in  
CC gene therapy. The method is useful for diagnosing and treating breast  
CC cancer.  
CC N.B. The sequence data for this patent did not form part of the printed  
CC specification, but was obtained in electronic format directly from WIPO  
CC at [ftp.wipo.int/pub/published\\_pct\\_sequences](http://wipo.int/pub/published_pct_sequences).  
XX  
SQ Sequence 9501 BP; 2816 A; 1872 C; 2078 G; 2735 T; 0 other;  
Query Match 80.0%; Score 16; DB 25; Length 9501;  
Best Local Similarity 100.0%; Pred. No. 8.6;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 2 CTTAGGAGGAAACAAGT 17  
Db 6556 CTTAGGAGGAAACAAGT 6571  
|||||  
RESULT 15  
ABK94929  
ID ABK94929 standard; cDNA; 9783 BP.  
XX  
XX ABK94929;  
AC  
XX  
XX 30-AUG-2002 (first entry)  
DT  
XX  
XX Human novel polynucleotide #40.  
DE  
XX  
XX Human; gene; ss; inflammatory condition; shock; sepsis; immune response;  
KW cancer; wound healing; central nervous system disease; haematopoiesis;  
KW peripheral nervous system disease; amyotrophic lateral sclerosis; tendon;  
KW myeloid cell disorder; lymphoid cell disorder; platelet disorder; bone;  
KW cartilage; ligament; nerve tissue; ulcer; osteoporosis; osteoarthritis;  
KW bone degenerative disorder; periodontal disease; reperfusion injury;  
KW lung fibrosis; liver fibrosis; autoimmune disorder; bacterial infection;  
KW allergic condition; thrombolytic; thrombosis; coagulation disorder;  
KW fungal infection.  
XX  
XX Homo sapiens.  
OS  
XX  
XX WO200244340-A2.  
PN  
XX  
XX 06-JUN-2002.  
PD  
XX  
XX 30-NOV-2001; 2001WO-UG47004.  
PF  
XX  
XX 30-NOV-2000; 2000US-0028952.  
PR  
XX  
XX (HXYSE-) HXYSEQ INC.  
PA  
XX  
XX Tang YT, Goodrich RW, Liu C, Zhou P, Asundi V, Wang J, Wang D;  
PI Yamazaki V, Ujwal ML, Drmanac RT;  
XX  
XX WPI; 2002-508509/54.  
DR  
XX  
XX P-PSDB; ABG66705.  
DR  
XX  
XX Novel nucleic acids and polypeptides for diagnosis, treatment of  
PT inflammatory, autoimmune, nervous system, myeloid or lymphoid cell  
PT disorders, cancer and promoting wound healing -  
XX  
XX Claim 1; Page 425-434; 672pp; English.  
XX  
XX The invention relates to human novel polynucleotides and associated  
CC polypeptides. The polynucleotides and polypeptides are useful for  
CC treating inflammatory conditions such as arthritis, nephritis, Crohn's  
CC disease, ischaemia-reperfusion injury, shock, sepsis, immune responses  
CC and cancer and for promoting wound healing. The sequences are used to  
CC induce the proliferation of neural cells and regeneration of nerve and  
CC brain tissue, and are useful for the treatment of central and peripheral  
CC nervous system diseases and neuropathies, such as Alzheimer's disease,

CC Parkinson's disease, Huntington's disease and amyotrophic lateral  
CC sclerosis. The sequences are involved in chemotactic or chemokinetic  
CC activity, regulation of haematopoiesis, treatment of myeloid or lymphoid  
CC cell disorders and platelet disorders such as thrombocytopenia,  
CC regeneration of bone, cartilage, tendon, ligament and/or nerve tissue  
CC growth, tissue repair, healing of burns, incisions, ulcers, treatment of  
CC osteoporosis, osteoarthritis, bone degenerative disorders and periodontal  
CC disease. The sequences of the invention are also useful for gut  
CC protection or regeneration and treatment of lung or liver fibrosis,  
CC reperfusion injury in various tissues, immune deficiencies and disorders  
CC including severe combined immunodeficiency (SCID), bacterial or fungal  
CC infections, autoimmune disorders e.g. multiple sclerosis and myasthenia  
CC gravis, allergic conditions such as asthma, thrombolytic or thrombosis  
CC and coagulation disorders. Sequences ABK94890-ABK94982 represent human  
CC novel polynucleotides of the invention.  
XX  
SQ Sequence 9783 BP; 2881 A; 1927 C; 2156 G; 2819 T; 0 other;  
Query Match 80.0%; Score 16; DB 24; Length 9783;  
Best Local Similarity 100.0%; Pred. No. 8.5;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 2 CTTAGGAGGAAACAAGT 17  
Db 6826 CTTAGGAGGAAACAAGT 6841  
|||||  
RESULT 16  
ABV22178  
ID ABV22178 standard; cDNA; 9785 BP.  
XX  
XX ABV22178;  
AC  
XX  
XX 13-SEP-2002 (first entry)  
DT  
XX  
XX Human prostate expression marker cDNA 22169.  
DE  
XX  
XX Human prostate cancer; cytostatic; carcinogen; pharmacodynamic marker;  
KW Human; prostate cancer; cytostatic; carcinogen; pharmacodynamic marker;  
KW pharmacogenomic marker; gene; ss.  
XX  
XX Homo sapiens.  
OS  
XX  
XX WO200160860-A2.  
PN  
XX  
XX 23-AUG-2001.  
PD  
XX  
XX 20-FEB-2001; 2001WO-US05171.  
PF  
XX  
XX 17-FEB-2000; 2000US-183319P.  
PR  
XX  
XX 16-MAR-2000; 2000US-189862P.  
PR  
XX  
XX 25-MAY-2000; 2000US-207454P.  
PR  
XX  
XX 09-JUN-2000; 2000US-211314P.  
PR  
XX  
XX 18-JUL-2000; 2000US-219007P.  
PR  
XX  
XX 13-DEC-2000; 2000US-255281P.  
PA  
XX  
XX (MILL-) MILLENNIUM PREDICTIVE MEDICINE INC.  
XX  
XX Schlegel R, Endege WO, Monahan JE;  
PI  
XX  
XX WPI; 2001-662795/76.  
DR  
XX  
XX Novel isolated nucleic acid molecule associated with cancerous state of  
PT prostate cells and correlating with presence of prostate cancer, useful  
PT for detecting presence of prostate cancer, stage of prostate cancer -  
XX  
XX Claim 1; Page 3813-3814; 11750pp; English.  
XX  
XX The invention relates to an isolated nucleic acid molecule (I) comprising  
CC a nucleotide sequence given in Tables 1-9 (ABV00010-ABV62213) of the  
CC specification or its complement. (I) is useful for:  
CC (a) assessing whether a patient is afflicted with prostate cancer;  
CC (b) monitoring the progression of prostate cancer in a patient;  
CC (c) assessing the efficacy of a test compound to inhibit prostate

CC cancer in a patient;  
CC (d) assessing the efficacy of a therapy for inhibiting prostate cancer  
CC in a patient;  
CC (e) selecting a composition for inhibiting prostate cancer in a patient;  
CC (f) assessing the prostate cell carcinogenic potential of a compound;  
CC (g) determining whether prostate cancer has metastasized in a patient;  
CC (h) assessing the aggressiveness or indolence of prostate cancer in a patient;  
CC (i) is also useful as a pharmacodynamic or pharmacogenomic marker.  
XX Sequence 9785 BP; 2870 A; 1936 C; 2160 G; 2819 T; 0 other;  
SQ  
Query Match 80.0%; Score 16; DB 23; Length 9785;  
Best Local Similarity 100.0%; Pred. No. 8.5;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 2 CTTAGGAGAACAACT 17  
DB 6835 CTTAGGAGAACAACT 6850  
RESULT 17  
ABV28017  
ID ABV28017 standard; cDNA; 9785 BP.  
XX  
AC ABV28017;  
XX  
DT 16-SEP-2002 (first entry)  
XX  
DE Human prostate expression marker cDNA 28008.  
XX  
KW Human; prostate cancer; cytostatic; carcinogen; pharmacodynamic marker;  
KW pharmacogenomic marker; gene; ss.  
XX  
OS Homo sapiens.  
XX  
PN W02C0160860-A2.  
XX  
PD 23-AUG-2001.  
XX  
PF 20-FEB-2001; 2001WO-US05171.  
XX  
PR 17-FEB-2000; 2000US-183119P.  
PR 16-MAR-2000; 2000US-189862P.  
PR 25-MAY-2000; 2000US-207454P.  
PR 09-JUN-2000; 2000US-271314P.  
PR 18-JUL-2000; 2000US-219007P.  
PR 13-DEC-2000; 2000US-255281P.  
XX  
PA (MILL-) MILLENNIUM PREDICTIVE MEDICINE INC.  
XX  
PI Schlegel R, Endege WO, Monahan JE;  
XX  
DR WPI; 2001-662795/76.  
XX  
PT Novel isolated nucleic acid molecule associated with cancerous state of  
PT prostate cells and correlating with presence of prostate cancer. useful  
PT for detecting presence of prostate cancer, stage of prostate cancer -  
XX  
PS Claim 1; Page 5783-5785; 11750DP; English.  
XX  
CC The invention relates to an isolated nucleic acid molecule (I) comprising  
CC a nucleotide sequence given in Tables 1-9 (ABV00010-ABV62213) of the  
CC specification or its complement. (I) is useful for:  
CC (a) assessing whether a patient is afflicted with prostate cancer;  
CC (b) monitoring the progression of prostate cancer in a patient;  
CC (c) assessing the efficacy of a test compound to inhibit prostate  
CC cancer in a patient;  
CC (d) assessing the efficacy of a therapy for inhibiting prostate cancer  
CC in a patient;  
CC (e) selecting a composition for inhibiting prostate cancer in a patient;  
CC (f) assessing the prostate cell carcinogenic potential of a compound;  
CC (g) determining whether prostate cancer has metastasized in a patient;

CC (h) assessing the aggressiveness or indolence of prostate cancer in a  
CC patient;  
CC (i) is also useful as a pharmacodynamic or pharmacogenomic marker.  
XX  
SQ Sequence 9785 BP; 2870 A; 1936 C; 2160 G; 2819 T; 0 other;  
SQ  
Query Match 80.0%; Score 16; DB 23; Length 9785;  
Best Local Similarity 100.0%; Pred. No. 8.5;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 2 CTTAGGAGAACAACT 17  
DB 6835 CTTAGGAGAACAACT 6850  
RESULT 18  
ABN85750/C  
ID ABN85750 standard; DNA; 180557 BP.  
XX  
AC ABN85750;  
XX  
DT 14-OCT-2002 (first entry)  
XX  
DE Human BAC clone RP11-334G22 SEQ ID NO 6.  
XX  
DE Human; Can 1; antiinfertility; gynaecological; infertility;  
KW premature ovarian failure; menopause; Sertoli Cell only syndrome;  
KW BAC clone RP11-334G22; GenBank reference AC007250; ds.  
XX  
OS Homo sapiens.  
XX  
PN US2002119929-A1.  
XX  
PD 29-AUG-2002.  
XX  
PF 02-NOV-2001; 2001US-0003806.  
XX  
PR 03-NOV-2000; 2000US-245872P.  
XX  
PA (BISH/) BISHOP C E.  
PA (AGOU/) AGOUTINIK A I.  
PA (ZHUQ/) ZHU Q.  
XX  
PI Bishop CE, Agoutinik AI, Zhu Q;  
XX  
DR WPI; 2002-618953/66.  
XX  
XX A nucleic acid molecule (I) encoding a Can 1 polypeptide used in  
PT treating infertility -  
XX  
PS Disclosure; Page -: 45pp; English.  
XX  
CC The invention relates to a nucleic acid molecule (I) encoding a Can 1  
CC polypeptide. The Can 1 nucleic acid molecule is used to diagnose or treat  
CC infertility or premature ovarian failure or Sertoli Cell only syndrome  
CC in a mammal. The present sequence is that of a human Can 1 encoding  
CC BAC clone RP11-334G22 of the invention.  
CC Note: The present sequence is not given in the printed specification but  
CC was obtained through the Genbank reference AC007250.  
XX  
SQ Sequence 180557 BP; 53238 A; 32016 C; 33894 G; 61409 T; 0 other;  
SQ  
Query Match 80.0%; Score 16; DB 24; Length 180557;  
Best Local Similarity 100.0%; Pred. No. 7.1;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 5 AGGAGAACAACTGCC 20  
DB 124080 AGGAGAACAACTGCC 124065  
RESULT 19  
AAH62689

ID AAH62689 standard; DNA; 305107 BP.  
 XX  
 AC AAH62689;  
 XX  
 DT 11-SEP-2001 (first entry)  
 XX  
 DE Shrimp white spot Bacilliform virus (WSBV) genomic sequence.  
 XX  
 KW Shrimp white spot Bacilliform virus; WSBV; diagnosis; viral infection;  
 KW antiviral agent; gene expression; antisense construct;  
 KW transgenic viral resistant shrimp; ds.  
 XX  
 OS White spot syndrome virus.  
 XX  
 PN WO200138351-A2.  
 XX  
 PD 31-MAY-2001.  
 XX  
 PF 08-NOV-2000; 2000WO-US28888.  
 XX  
 PR 24-NOV-1999; 99CN-0124717.  
 XX  
 PA (PENY-) PE CORP NY.  
 PA (THIR-) THIRD INST OCEANOGRAPHY STATE OCEANI C A.  
 PA (SINO-) SINOGENOMAX CO LTD.  
 XX  
 PI Xu X, Yang F, He J, Pham L, He M, Ye Y, Shen Y, Kodira C;  
 XX  
 DR WPI; 2001-355877/37.  
 XX  
 PT Primary nucleotide sequence of the shrimp white spot Bacilliform virus  
 PT (WSBV), useful for producing viral polypeptides that can be used to  
 PT screen for agents that are useful for treating WSBV infection -  
 XX  
 PS Disclosure; Figure 2; 626pp; English.  
 XX  
 CC The invention provides the primary nucleotide sequence of the WSBV genome  
 CC (AAH62689), predicted transcript sequences (AAH62689-AAH62839) and  
 CC encoded proteins (AAG64910-AAG65051) and oligonucleotide sequences  
 CC (AAH62840-63160) suitable for use as primers or probes. The nucleic acid  
 CC molecules and proteins of the invention are useful for diagnosis and  
 CC monitoring viral infection, in screens for antiviral agents and for  
 CC monitoring viral gene expression or activity during a treatment regimen.  
 CC The nucleic acid molecules are also useful as antisense constructs to  
 CC control viral gene expression in infected cells and tissues and to create  
 CC transgenic viral resistant shrimp.  
 XX  
 SQ Sequence 305107 BP; 92042 A; 62482 C; 62635 G; 87948 T; 0 other;  
 XX  
 QY  
 Db 5 AGGAGGAACAAGTCCC 20  
 155349 AGGAGGAACAAGTCCC 155364  
 XX  
 RESULT 20  
 AAC20071  
 ID AAC20071 standard; cDNA; 123 BP.  
 XX  
 AC AAC20071;  
 XX  
 DT 06-OCT-2000 (first entry)  
 XX  
 DE Human secreted protein 5' EST, SEQ ID NO: 24146.  
 XX  
 KW Human; 5' EST; expressed sequence tag; secreted protein; cDNA isolation;  
 KW gene therapy; chromosome mapping; ss.  
 XX  
 OS Homo sapiens.  
 XX

PN EP1033401-A2.  
 XX  
 PD 06-SEP-2000.  
 XX  
 PF 21-FEB-2000; 2000EP-0200610.  
 XX  
 PR 26-FEB-1999; 99US-0122487.  
 XX  
 PA (GEST) GENSET.  
 XX  
 PI Dumas Milne Edwards J, Duclert A, Giordano J;  
 XX  
 DR WPI; 2000-500381/45.  
 XX  
 PT New nucleic acid that is a 5' expressed sequence tag (5' EST) for  
 PT obtaining cDNAs and genomic DNAs that correspond to 5'ESTs and for  
 PT diagnostic, forensic, gene therapy and chromosome mapping procedures -  
 XX  
 PS Claim 1; SEQ ID 24146; 71pp + CD-ROM; English.  
 XX  
 CC The present sequence is one of a large number of 5' ESTs derived from  
 CC cDNAs encoding secreted proteins. No ORF has yet been conclusively  
 CC identified within the present sequence. The 5' ESTs were prepared from  
 CC total human RNAs or polyA+ RNAs derived from 30 different tissues. EST  
 CC sequences usually correspond mainly to the 3' untranslated region (UTR)  
 CC of the mRNA because they are often obtained from oligo-dT primed cDNA  
 CC libraries. Such ESTs are not well suited for isolating cDNA sequences  
 CC derived from the 5' ends of mRNAs and even in those cases where longer  
 CC cDNA sequences have been obtained, the full 5' UTR is rarely included.  
 CC 5' ESTs are derived from mRNAs with intact 5' ends and can therefore be  
 CC used to obtain full length cDNAs and genomic DNAs. 5' ESTs are also used  
 CC in diagnostic, forensic, gene therapy and chromosome mapping procedures.  
 CC They are used to obtain upstream regulatory sequences and to design  
 CC expression and secretion vectors.  
 XX  
 SQ Sequence 123 BP; 48 A; 15 C; 22 G; 33 T; 5 other;  
 XX  
 QY  
 Db 3 TTAGAGGACCAAGT 17  
 19 TTAGAGGACCAAGT 33  
 XX  
 RESULT 21  
 ABK99951/C  
 ID ABK99951 standard; DNA; 762 BP.  
 XX  
 AC ABK99951;  
 XX  
 DT 21-OCT-2002 (first entry)  
 XX  
 DE DNA encoding human secreted protein SCEP-34.  
 XX  
 KW Secreted protein; SCEP; human; cell proliferative disorder; cancer;  
 KW keratosis; arteriosclerosis; atherosclerosis; cirrhosis; hepatitis;  
 KW psoriasis; autoimmune disorder; inflammatory disorder; AIDS; arthritis;  
 KW acquired immunodeficiency syndrome; adult respiratory distress syndrome;  
 KW Addison's disease; allergy; asthma; osteoporosis; autoimmune thyroiditis;  
 KW Crohn's disease; dermatitis; diabetes; Graves' disease; haemodialysis;  
 KW glomerulonephritis; scleroderma; systemic lupus erythematosus; uveitis;  
 KW systemic sclerosis; ulcerative colitis; infection; trauma; Pick disease;  
 KW cardiovascular disorder; congestive heart failure; angina; epilepsy;  
 KW hypertensive heart disease; neurological disorder; Alzheimer's disease;  
 KW Parkinson's disease; amyotrophic lateral sclerosis; stroke; dementia;  
 KW Huntington's disease; multiple sclerosis; neuromuscular disorder;  
 KW metabolic disorder; endocrine disorder; toxic myopathy; mental disorder;  
 KW schizophrenia; developmental disorder; anaemia; epilepsy;  
 KW hypothyroidism; glaucoma; sensorineural hearing loss; cataract;  
 KW transgenic animal; gene; ds.  
 XX

OS Homo sapiens.  
 XX WO200248337-A2.  
 XX  
 PD 20-JUN-2002.  
 XX  
 PF 12-DEC-2001; 2001WO-US48517.  
 XX  
 PR 13-DEC-2000; 2000US-255639P.  
 PR 21-DEC-2000; 2000US-257852P.  
 PR 05-JAN-2001; 2001US-260105P.  
 PR 18-JAN-2001; 2001US-262932P.  
 PR 18-JAN-2001; 2001US-263096P.  
 PR 19-JAN-2001; 2001US-263090P.  
 PR 02-FEB-2001; 2001US-265926P.  
 XX  
 PA (INCY-) INCYTE GENOMICS INC.  
 XX  
 PI Griffin JA, Yao MG, Duggan BM, Yue H, Ding L, Lal PG, Lee EA,  
 PI Ramkumar J, Thangavelu K, Xu Y, Lee S, Tang YT, Nguyen DB,  
 PI Warren BA, Honchell CD, Gietzen KJ, Baughn MR, Gandhi AR,  
 PI Arvizu C, Walla NK, Lu Y, Elliott VM, Lu DAM, Hafalia AJA;  
 PI Azimzai Y, Khan FA, Tran UK;  
 XX  
 DR WPI; 2002-583509/62.  
 DR P-PSDB; ABG69654.  
 XX  
 PT Novel human secreted proteins and polynucleotides for diagnosing,  
 PT preventing or treating cell proliferative, autoimmune/inflammatory,  
 PT cardiovascular, neurological and developmental disorders -  
 XX  
 PS Claim 5; Page 221; 234pp; English.  
 XX  
 CC The invention describes an isolated polypeptide chosen from secreted  
 CC proteins (I), SECP 1-54. (I) and the polynucleotide encoding it (II) are  
 CC useful for screening a compound for effectiveness as an agonist or  
 CC antagonist of (I) or compound that alters expression of (II). (I), the  
 CC identified agonist and antagonist are useful for treating a disease or  
 CC condition associated with altered expression of functional SECP in a patient.  
 CC An antibody specific to (I) is useful for detecting the presence of (I),  
 CC purifying (I) from a sample and for diagnosing a condition or disease  
 CC associated with expression of SECP in a subject or in a biological  
 CC sample. (I) and (II) and modulators of (I) are useful for diagnosis,  
 CC treatment and prevention of cell proliferative disorders (e.g. cancer,  
 CC keratosis, arteriosclerosis, atherosclerosis, cirrhosis, hepatitis and  
 CC poriasis), autoimmune/inflammatory disorders (e.g. acquired  
 CC immunodeficiency syndrome (AIDS), adult respiratory distress syndrome,  
 CC Addison's disease, allergies, asthma, osteoporosis, autoimmune  
 CC thyroiditis, Crohn's disease, dermatitis, diabetes, Graves' disease,  
 CC glomerulonephritis, arthritis, scleroderma, systemic lupus erythematosus,  
 CC systemic sclerosis, ulcerative colitis, haemodialysis, uveitis; viral,  
 CC bacterial, fungal, parasitic, protozoal, helminthic infections and  
 CC trauma), cardiovascular disorders (e.g. congestive heart failure, angina,  
 CC hypertensive heart disease), neurological disorders (e.g. Alzheimer's and  
 CC Pick disease, Parkinson's disease, amyotrophic lateral sclerosis, epilepsy,  
 CC stroke, Huntington's disease, multiple sclerosis, dementia, neuromuscular  
 CC disorders, metabolic, endocrine and toxic myopathies, mental disorders,  
 CC schizophrenic disorders, and developmental disorders (e.g. anaemia,  
 CC epilepsy, hypothyroidism, glaucoma, sensorineural hearing loss and  
 CC cataract). (II) is useful for creating transgenic animals to model human  
 CC disease and to detect and quantify gene expression in biopsied tissues in  
 CC which expression of SECP is correlated with disease. This sequence  
 CC encodes a human secreted protein (SECP).  
 XX  
 SQ Sequence 762 BP; 174 A; 210 C; 164 G; 214 T; 0 other;

Query Match 75.0%; Score 15; DB 24; Length 762;  
 Best Local Similarity 100.0%; Pred. No. 36;  
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 5 AGGAGGAACAGTCC 19  
 |||||  
 DB 555 AGGAGGAACAGTCC 541

RESULT 22  
 ABZ66827  
 ID ABZ66827 standard; DNA; 1691 BP.  
 XX  
 AC ABZ66827;  
 XX  
 DT 24-MAR-2003 (first entry)  
 XX  
 DE Arabidopsis thaliana polynucleotide SEQ ID NO 23.  
 XX  
 KW Arabidopsis thaliana; herbicidal; plant; growth regulator; transgenic;  
 KW herbicide-resistant; gene; ds.  
 XX  
 OS Arabidopsis thaliana.  
 XX  
 FN WO200266660-A2.  
 PD  
 PD 29-AUG-2002.  
 XX  
 PF 13-FEB-2002; 2002WO-BP01466.  
 XX  
 PR 16-FEB-2001; 2001DE-1007843.  
 PR 23-MAY-2001; 2001DE-1025537.  
 XX  
 PA (META-) METANOMICS GMBH & CO KGAA.  
 XX  
 PI Plesch G, Blau A, Daeschner K, Klein M;  
 XX  
 DR WPI; 2002-674953/72.  
 XX  
 PT Identifying herbicides and plant growth regulators, from ability to  
 PT inhibit specific genes; also use of these genes to prepare  
 PT herbicide-resistant transgenic plants -  
 XX  
 PS Disclosure; Page 154-155; 224pp; German.  
 XX  
 CC The invention relates to identifying compounds (A) with herbicidal  
 CC activity from their ability to reduce or block expression or activity of  
 CC gene products of specific nucleic acid sequences (I) or amino acid  
 CC sequences (II) encoded by (I). (A) and antagonists of proteins encoded  
 CC by (I) are useful as herbicides and plant growth regulators.  
 CC Overexpression of (I) in plants is used to produce herbicide-resistant  
 CC plants and (I) may also be mutated to identify altered sequences that  
 CC encode proteins resistant to herbicides. The present sequence is that of  
 CC a polynucleotide of the invention.  
 XX  
 SQ Sequence 1691 BP; 483 A; 309 C; 434 G; 465 T; 0 other;

Query Match 75.0%; Score 15; DB 24; Length 1691;  
 Best Local Similarity 100.0%; Pred. No. 34;  
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 3 TTAGGAGGAACAGT 17  
 |||||  
 DB 830 TTAGGAGGAACAGT 844

RESULT 23  
 ABL29548  
 ID ABL29548 standard; DNA; 17255 BP.  
 XX  
 AC ABL29548;  
 XX  
 DT 26-MAR-2002 (first entry)  
 XX  
 DE Drosophila melanogaster genomic polynucleotide SEQ ID NO 40117.  
 XX  
 KW Drosophila; developmental biology; cell signalling; insecticide;  
 KW pharmaceutical; gene; ds.  
 XX  
 OS Drosophila melanogaster.

```

XX PN WO200171042-A2.
XX PD
XX 27-SEP-2001.
XX PF 23-MAR-2001; 2001WO-US09231.
XX PR 23-MAR-2000; 2000US-191637P.
XX PR 11-JUL-2000; 2000US-0614150.
XX PA (PEKE ) PE CORP NY.
XX PI Venter JC, Adams M, Li PWD, Myers EW;
XX WP1; 2001-656860/75.
XX
XX PT New isolated nucleic acid detection reagent for detecting 1000 or more
XX PT genes from Drosophila and for elucidating cell signalling and cell-cell
XX PT interactions -
XX
XX PS Claim 1; SEQ ID NO 40117; 21pp + Sequence listing; English.
XX
XX CC The invention relates to an isolated nucleic acid detection reagent
XX CC capable of detecting 1000 or more genes from Drosophila. The invention is
XX CC useful in developmental biology and in elucidating cell signalling and
XX CC cell-cell interactions in higher eukaryotes for the development of
XX CC insecticides, therapeutics and pharmaceutical drugs. The invention
XX CC discloses genomic DNA sequences (AB16176-ABL30511), expressed DNA
XX CC sequences (AB101840-ABL16175) and the encoded proteins
XX CC (ABB57737-ABB72072).
XX CC The sequence data for this patent did not form part of the printed
XX CC specification, but was obtained in electronic format directly from WIPO
XX CC at ftp.wipo.int/pub/published_pct_sequences.
XX
XX SO Sequence 17255 BP; 5336 A; 3353 C; 3445 G; 5121 T; 0 other;

OY 5 AGGAGGAACAAGTCC 19
Db 10781 AGGAGGAACAAGTCC 10795

RESULT 24
ABQ76896
ID ABQ76896 standard; DNA; 53226 BP.
XX
XX AC ABQ76896;
XX
XX DT 13-MAR-2003 (first entry)
XX
DE Human G-protein coupled receptor DNA SEQ ID 3.
XX
XX G-protein coupled receptor; secretin receptor subfamily; human; SNP;
XX GPCR; protease; Parkinson's disease; gene; chromosome X;
XX single nucleotide polymorphism; ds.
XX
XX Homo sapiens.
XX
XX Key Location/Qualifiers
XX CDS 3000..50651
XX /tag= a
XX /product= "GPCR"
XX /note="this coding sequence is interrupted by
XX 13 introns"
XX replace (1746..c)
XX /tag= b
XX /note= "SNP, single nucleotide polymorphism"
XX replace (1755..g)
XX /tag= c
XX /note= "SNP, single nucleotide polymorphism"
XX
XX variation
XX
XX variation
XX
XX variation
XX

```

FT	variation	replace (1961,g)	/*tag= d
FT		/note= "SNP, single nucleotide polymorphism"	
FT	exon	3000..3088	/*tag= e
FT		/number= 1	
FT	intron	3089..3874	/*tag= f
FT		/number= 1	
FT	exon	3875..4038	/*tag= g
FT		/number= 2	
FT	intron	4039..6037	/*tag= h
FT		/number= 2	
FT	variation	replace (5411,g)	/*tag= i
FT		/note= "SNP, single nucleotide polymorphism"	
FT	variation	replace (5760,a)	/*tag= j
FT		/note= "SNP, single nucleotide polymorphism"	
FT	exon	6038..6170	/*tag= k
FT		/number= 3	
FT	intron	6171..8059	/*tag= l
FT		/number= 3	
FT	exon	8060..8178	/*tag= m
FT		/number= 4	
FT	intron	8179..15910	/*tag= n
FT		/number= 4	
FT	variation	replace (11390,c)	/*tag= o
FT		/note= "SNP, single nucleotide polymorphism"	
FT	variation	replace (11390,t)	/*tag= p
FT		/note= "SNP, single nucleotide polymorphism"	
FT	exon	15911..16127	/*tag= q
FT		/number= 5	
FT	intron	16128..17484	/*tag= r
FT		/number= 5	
FT	variation	replace (16988,a)	/*tag= s
FT		/note= "SNP, single nucleotide polymorphism"	
FT	exon	17485..17647	/*tag= t
FT		/number= 6	
FT	intron	17648..32332	/*tag= u
FT		/number= 6	
FT	variation	replace (18361,g)	/*tag= v
FT		/note= "SNP, single nucleotide polymorphism"	
FT	variation	replace (19769,c)	/*tag= w
FT		/note= "SNP, single nucleotide polymorphism"	
FT	variation	replace (22910,t)	/*tag= x
FT		/note= "SNP, single nucleotide polymorphism"	
FT	variation	replace (22935,t)	/*tag= y
FT		/note= "SNP, single nucleotide polymorphism"	
FT	variation	replace (24206,g)	/*tag= z
FT		/note= "SNP, single nucleotide polymorphism"	
FT	variation	replace (24774,a)	/*tag= aa
FT		/note= "SNP, single nucleotide polymorphism"	
FT	variation	replace (24869,c)	/*tag= ab

```
FT      /*tag= ab
FT      /note= "SNP, single nucleotide polymorphism"
FT      replace (25768,a)
FT      /*tag= ac
FT      /note= "SNP, single nucleotide polymorphism"
FT      replace (26697,c)
FT      /*tag= ad
FT      /note= "SNP, single nucleotide polymorphism"
FT      replace (26697,c)
FT      /*tag= ae
FT      /note= "SNP, single nucleotide polymorphism"
FT      replace (28359,c)
FT      /*tag= af
FT      /note= "SNP, single nucleotide polymorphism"
FT      replace (28470..28471,t)
FT      /*tag= ag
FT      /note= "a single nucleotide polymorphism (SNP) can
FT      result in a deletion at this position"
FT      replace (29781,g)
FT      /*tag= ah
FT      /note= "SNP, single nucleotide polymorphism"
FT      replace (30182,a)
FT      /*tag= ai
FT      /note= "SNP, single nucleotide polymorphism"
FT      replace (31772,t)
FT      /*tag= aj
FT      /note= "SNP, single nucleotide polymorphism"
FT      replace (31936,g)
FT      /*tag= ak
FT      /note= "SNP, single nucleotide polymorphism"
FT      /*tag= al
FT      /number= 7
FT      32468..36825
FT      /*tag= am
FT      /number= 7
FT      36826..36948
FT      /*tag= an
FT      /product= 8
FT      36949..38129
FT      /*tag= ao
FT      /number= 8
FT      38130..38175
FT      /*tag= ap
FT      /number= 9
FT      38176..39984
FT      /*tag= aq
FT      /number= 9
FT      39985..40088
FT      /*tag= ar
FT      /number= 10
FT      40089..42455
FT      /*tag= as
FT      /number= 10
FT      42456..42577
FT      /*tag= at
FT      /number= 11
FT      42578..44422
FT      /*tag= au
FT      /number= 11
FT      replace (42767..42767,c)
FT      /*tag= av
FT      /note= "a single nucleotide polymorphism (SNP) can
FT      result in a deletion at this position"
FT      44423..44691
FT      /*tag= aw
FT      /number= 12
FT      44692..47818
FT      /*tag= ax
FT      /number= 12
FT      47819..47897
FT      /*tag= ay
FT      /number= 13
```

```
FT      intron
FT      47898..50266
FT      /*tag= az
FT      /number= 13
FT      replace (48839,c)
FT      /*tag= ba
FT      /note= "SNP, single nucleotide polymorphism"
FT      50267..50651
FT      /*tag= bb
FT      /number= 14
FT      replace (52265,g)
FT      /*tag= bc
FT      /note= "SNP, single nucleotide polymorphism"
FT      US2002142951-A1.
FT      PN
FT      03-OCT-2002.
FT      PD
FT      28-MAR-2001; 2001US-0818264.
FT      PR
FT      28-MAR-2001; 2001US-0818264.
FT      XX
FT      28-MAR-2001; 2001US-0818264.
FT      PR
FT      (WEBS/) WEBSTER M.
FT      PA (BEAS/) BEASLEY E. M.
FT      PA (KETC/) KETCHUM K. A.
FT      PA (DFRA/) DI FRANCESCO V.
FT      XX
FT      PI Webster M., Beasley EM, Ketchum KA, Di Francesco V;
FT      XX
FT      Query Match 75.0%; Score 15; DB 25; Length 53226;
FT      Best Local Similarity 100.0%; Pred. No. 27;
FT      Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
FT      Oy 2 CTTAGAGAACCAAG 16
FT      Db 35513 CTTAGAGAACCAAG 35527
FT      ID AAQ76831
FT      XX AAQ76831;
FT      AC
FT      XX 25-MAR-2003 (updated)
FT      DT 23-SEP-1994 (first entry)
FT      XX
FT      DE Human genome fragment.
FT      XX
FT      KW Brain; placenta; bone marrow; genetic analysis; gene mapping;
FT      KW detection; homology; human; adrenal tissue; ds.
FT      XX
FT      OS Homo sapiens.
FT      XX
FT      PN W09401548-A2.
FT      PN
FT      PD 20-JAN-1994.
FT      PD
FT      XX 13-JUL-1993; 93MO-GB01467.
FT      XX
FT      PR 13-JUL-1992; 92GB-0014857.
FT      XX
FT      PA (MED-) MEDICAL RES COUNCIL.
FT      XX
FT      PI Gross J, Hadfield KM, Howells D, Kelly M, Shaw D;
FT      PI Sidson DR, Starkey M;
FT      DR WPI; 1994-035056/04.
FT      XX
FT      New nucleic acid fragment encoding gene products - can be used
FT      for genetic analysis and mapping
FT      PS Claim 1; Page 228; 616bp; English.
```

XX Human nucleic acid fragments, isolated from brain, adrenal tissue,  
CC the placenta or bone marrow comprise any of: (A) a sequence  
CC selected from (AA076401-Q77613), (B) an allelic variation of a  
CC sequence as described in (A), or (C) a sequence complementary  
CC to (A) or (B).  
CC Preferred sequences exhibit no more than 90% homology to a human  
CC sequence known per se.  
CC (Updated on 25-MAR-2003 to correct PN field.)  
XX  
SQ Sequence 125 BP; 29 A; 31 C; 39 G; 25 T; 1 other;  
Query Match 70.0%; Score 14; DB 15; Length 125;  
Best Local Similarity 100.0%; Pred. No. 1.4e+02;  
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 5 AGGAGGACCAAGTC 18  
DB 106 AGGAGGACCAAGTC 119  
RESULT 26  
ABX30187  
ID ABX30187 standard; cDNA; 273 BP.  
XX  
AC ABX30187;  
XX  
DT 11-FEB-2003 (first entry)  
XX  
DE Human GDP-mannose 4,6-dehydratase (GM4,6D) DNA #12244.  
XX  
KW Human; GDP-mannose 4,6-dehydratase; GM4,6D; gene; ss; inflammation;  
KW cellular fucosylation; glycoconjugate fucosylation; transplant rejection;  
KW arthritis; asthma; sepsis; reperfusion injury; stroke; infection;  
KW complex carbohydrate; gene replacement therapy; immunosuppressive;  
KW antiinflammatory; antiarthritic; antibacterial; cerebroprotective;  
KW antiasthmatic; vasotropic.  
XX  
OS Homo sapiens.  
XX  
PN US2002110548-A1.  
XX  
PD 15-AUG-2002.  
XX  
PF 11-JUN-2001; 2001US-0878574.  
XX  
PR 22-NOV-1996; 96US-0753233.  
PR 03-DEC-1997; 97US-0984246.  
PR 09-SEP-1998; 98US-0149674.  
PR 14-JUN-1999; 99US-0333177.  
XX  
PA (GENY) GENETICS INST INC.  
XX  
PI Sullivan F, Kriz R, Kumar R;  
XX  
DR WPI; 2003-066673/06.  
XX  
PT New composition comprising GDP-mannose 4,6-dehydratase (GM4,6D)  
PT peptide, for manufacturing complex carbohydrates, or as targets for  
PT screening GM4,6D antagonists for treating e.g. arthritis, or transplant  
PT rejection -  
XX  
PS Disclosure; SEQ ID NO 12246; 6pp; English.  
XX  
CC The invention relates to a composition comprising a human GDP-mannose  
CC 4,6-dehydratase (GM4,6D) peptide. The peptide is useful for identifying  
CC GM4,6D inhibitors. GM4,6D inhibitors are useful for reducing inflammation  
CC in a mammalian subject and for treating or ameliorating diseases affected  
CC by the level of cellular fucosylation or diseases affected by the  
CC fucosylation of glycoconjugates. These diseases include arthritis,  
CC transplant rejection, asthma, sepsis, reperfusion injury, stroke or  
CC infection. The GM4,6D peptide or a polynucleotide encoding it is also  
CC useful for manufacturing complex carbohydrates and as targets for

CC screening small molecule antagonists of the activity of the enzyme. The  
CC polynucleotide is useful in developing an assay for defects in the  
CC enzyme, as well as in gene replacement therapy. Sequences  
CC ABX17943-ABX17944 and ABX17947-ABX33716 represent DNA molecules encoding  
CC human GM4,6D peptides of the invention.  
CC Note: The sequence data for this patent did not form part of the printed  
CC specification but was obtained in electronic format directly from USPTO  
CC at [seqdata.uspto.gov/sequence.html](http://seqdata.uspto.gov/sequence.html).  
XX  
SQ Sequence 273 BP; 81 A; 57 C; 58 G; 77 T; 0 other;  
Query Match 70.0%; Score 14; DB 25; Length 273;  
Best Local Similarity 100.0%; Pred. No. 1.3e+02;  
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 2 CTTAGGAGGACCA 15  
DB 46 CTTAGGAGGACCA 59  
RESULT 27  
AAK98399/C  
ID AAK98399 standard; cDNA; 300 BP.  
XX  
AC AAK98399;  
XX  
DT 24-SEP-1999 (first entry)  
XX  
DE Human cancer cell derived cDNA #125.  
XX  
KW Cancer; human; colon; breast; lung; transmembrane receptor; ATPase;  
KW integral membrane protein; aspartyl protease; GATA family; wnt family;  
KW transcription factor; G-protein alpha subunit; protein phosphatase;  
KW phospholipase binding protein; diacylglycerol binding protein; trypsin;  
KW protein kinase; tyrosine phosphatase; developmental signalling protein;  
KW Wnt/PCP/Wnt domain; therapy; forensic; genetic mapping; diagnostic;  
KW detection; treatment; cervical; melanoma; colorectal adenocarcinoma;  
KW Wilms' tumour; retinoblastoma; sarcoma; myosarcoma; lung carcinoma;  
KW leukemia; lymphoma; dysplasia; hyperplasia; endometrium; adrenal;  
KW prostate; ss.  
XX  
OS Homo sapiens.  
XX  
PN WO9333982-A2.  
XX  
PD 08-JUL-1999.  
XX  
PF 22-DEC-1998; 98WO-US27610.  
XX  
PR 21-DEC-1998; 98US-0217471.  
PR 23-DEC-1997; 97US-0068755.  
PR 03-APR-1998; 98US-0080664.  
PR 21-OCT-1998; 98US-0105234.  
PR 27-OCT-1998; 98US-0105877.  
XX  
PA (CHIR) CHIRON CORP.  
XX  
PI (HYSE-) HYSEQ INC.  
XX  
DR Ckvenjakov R, Dickson M, Drmanac R, Drmanac S;  
XX  
PI Escobedo J, Garcia PD, Garcia V, Giese K, Imis MA;  
XX  
PI Jones LW, Kassam A, Kennedy GC, Kita D, Labat I;  
XX  
PI Lamson G, Leskowitz D, Pot D, Randazzo F, Reinhard C;  
XX  
PI Strache-Crain B, Sudduth-Klinger J, Williams LT;  
XX  
DR WPI; 1999-430243/36.  
XX  
PT New isolated human polynucleotides  
XX  
PS Claim 1; Page 348; 59pp; English.  
XX  
CC This invention describes novel isolated human polynucleotides obtained  
CC by screening for differential expression in colon cancer, breast cancer  
CC and lung cancer cell lines. The polynucleotides of the invention are

represented in AAX98275-X99118 and encode polypeptides of protein families selected from 4 transmembrane segments integral membrane proteins, 7 transmembrane receptors, ATPases associated with various cellular activities (AAA), eukaryotic aspartyl proteases, GATA family of transcription factors, G-protein alpha subunit, photolabile or diacylglycerol binding proteins, protein kinase, protein phosphatase 2C, protein tyrosine phosphatase, trypsin, wnt family of developmental signaling proteins and WW/rps5/WWP domain containing proteins. The encoded polypeptides also have a functional domain selected from Ank repeat, basic region plus leucine zipper transcription factors, bromodomain, EF-hand, SH3 domain, WD domain/G-beta repeats, zinc finger (CCH2 type), zinc finger (CCHC class), and zinc-binding metalloprotease domain. The polynucleotides encode polypeptides with similarity to known protein families and are predicted to have similar properties. The novel polynucleotides can be used to develop products for use as therapeutic agents and in forensics, genetic analysis, mapping and diagnostic applications. In particular, the product can be used for the detection and management of cancers. They can be used for treating e.g. cervical cancers, melanomas, colorectal adenocarcinomas, Wilms' tumour, sarcomas, retinoblastoma, myosarcomas, lung carcinomas, leukemias, such as chronic myelogenous leukemia, promyelocytic leukemia, monocytic leukemia, and myeloid leukemia, and lymphomas such as histiocytic lymphoma, anaplastic hereditary ectodermal dysplasia, congenital alveolar dysplasia, epithelial dysplasia of the cervix, fibrous dysplasia of bone, and mammary dysplasia, hyperplasias, e.g. endometrial, adrenal, breast, prostate or thyroid hyperplasias or pseudoepitheliomatous hyperplasia of the skin.

Sequence 300 BP, 81 A; 35 C; 63 G; 121 T; 0 other;

Query Match 70.0%; Score 14; DB 20; Length 300;  
Best Local Similarity 100.0%; Pred. No. 1.3e+02;  
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

1 CCTTAGGAGGAACA 14  
|||||  
230 CCTTAGGAGGAACA 217

RESULT 28  
ABV06501  
ID ABV06501 standard; cDNA; 344 BP.

AC ABV06501;

DT 13-SEP-2002 (first entry)

DE Human prostate expression marker cDNA 6492.

KW Human; prostate cancer; cytosolic; carcinogen; pharmacodynamic marker;

KM pharmacogenomic marker; gene; ss.

OS Homo sapiens.

PN MO200160860-A2.

PD 23-AUG-2001.

PF 20-FEB-2001; 2001WO-US05171.

PR 17-FEB-2000; 2000US-183319P.

PR 16-MAR-2000; 2000US-189862P.

PR 25-MAY-2000; 2000US-207454P.

PR 09-JUN-2000; 2000US-211314P.

PR 18-JUL-2000; 2000US-219007P.

PR 13-DEC-2000; 2000US-255281P.

PA (MILL-) MILLENNIUM PREDICTIVE MEDICINE INC.

PI Schlegel R, Endege WO, Nonahan JE;

XX WPI; 2001-662795/76.

PT Novel isolated nucleic acid molecule associated with cancerous state of prostate cells and correlating with presence of prostate cancer, useful for detecting presence of prostate cancer, stage of prostate cancer -

PS Claim 1; Page 1063; 11750pp; English.

CC The invention relates to an isolated nucleic acid molecule (I) comprising a nucleotide sequence given in Tables 1-9 (ABV00010-ABV62213) of the

CC specification or its complement. (I) is useful for:

CC (a) assessing whether a patient is afflicted with prostate cancer;

CC (b) monitoring the progression of prostate cancer in a patient;

CC (c) assessing the efficacy of a test compound to inhibit prostate cancer in a patient;

CC (d) assessing the efficacy of a therapy for inhibiting prostate cancer in a patient;

CC (e) selecting a composition for inhibiting prostate cancer in a patient;

CC (f) assessing the prostate cell carcinogenic potential of a compound;

CC (g) determining whether prostate cancer has metastasized in a patient;

CC (h) assessing the aggressiveness or indolence of prostate cancer in a patient;

CC (I) is also useful as a pharmacodynamic or pharmacogenomic marker.

Sequence 344 BP, 113 A; 52 C; 82 G; 94 T; 3 other;  
Query Match 70.0%; Score 14; DB 23; Length 344;  
Best Local Similarity 100.0%; Pred. No. 1.3e+02;  
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

5 AGAGGAGAACAGTC 18  
|||||  
100 AGAGGAGAACAGTC 113

RESULT 29  
AAC03189  
ID AAC03189 standard; cDNA; 357 BP.

AC AAC03189;

DT 06-OCT-2000 (first entry)

DE Human secreted protein 5' EST, SEQ ID NO: 3187.

KW Human; 5' EST; expressed sequence tag; secreted protein; cDNA isolation;

KM gene therapy; chromosome mapping; ss.

OS Homo sapiens.

PN EP1033401-A2.

PD 06-SEP-2000.

PF 21-FEB-2000; 2000EP-0200610.

PR 26-FEB-1999; 99US-0122487.

PR (GEST ) GENSET.

PI Dumas Milne Edwards J, Duclert A, Giordano J;

XX WPI; 2000-500381/45.

XX P-PSDB; AAG03189.

XX New nucleic acid that is a 5' expressed sequence tag (5' EST) for

XX obtaining cDNAs and genomic DNAs that correspond to 5' ESTs and for

XX diagnostic, forensic, gene therapy and chromosome mapping procedures -

XX Claim 1; SEQ ID 3187; 71pp + CD-ROM; English.

XX The present sequence is one of a large number of 5' ESTs derived from

XX mRNAs encoding secreted proteins. An ORF has been identified within the

XX sequence. The 5' ESTs were prepared from total human RNAs or polyA+ RNAs

XX derived from 30 different tissues. EST sequences usually correspond



CC mainly to the 3' untranslated region (UTR) of the mRNA because they are  
 CC often obtained from oligo-dT primed cDNA libraries. Such ESTs are not  
 CC well suited for isolating cDNA sequences derived from the 5' ends of  
 CC mRNAs and even in those cases where longer cDNA sequences have been  
 CC obtained, the full 5' UTR is rarely included. 5' ESTs are derived from  
 CC mRNAs with intact 5' ends and can therefore be used to obtain full length  
 CC cDNAs and genomic DNA. 5' ESTs are also used in diagnostic, forensic,  
 CC gene therapy and chromosome mapping procedures. They are used to obtain  
 CC upstream regulatory sequences and to design expression and secretion  
 CC vectors.

XX  
 SQ Sequence 357 BP, 112 A, 74 C, 78 G, 92 T, 1 other;

Query Match 70.0%; Score 14; DB 21; Length 357;  
 Best Local Similarity 100.0%; Pred. No. 1.3e+02;

Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 4 TAGGAGGACCACT 17  
 DB 223 TAGGAGGACCACT 236

RESULT 30  
 AAS38136/c  
 ID AAS38136 standard; cDNA; 408 BP.

XX  
 AC AAS38136;

XX 17-DEC-2001 (first entry)

XX Novel human diagnostic and therapeutic gene #1194.

XX Human; cancer; breast; lung; colon; prostate; cytostatic; diagnostic; ss.

XX Homo sapiens.

XX WO200166753-A2.

XX 13-SEP-2001.

XX 09-MAR-2001; 2001MO-US07787.

XX 09-MAR-2000; 2000US-0188609.

XX (CHIR ) CHIRON CORP.

XX (HYSE ) HYSEO INC.

XX Williams LT, Escobedo J, Innis MA, Garcia PD, Sudduth-Klinger J;  
 PI Reinhard C, Randazzo F, Kennedy GC, Pot D, Kassam A, Lamson G;  
 PI Drmanac R, Crkvenjakov R, Dickson M, Drmanac S, Labat I;  
 PI Leshkowitz D, Kita D, Garcia V, Jones WL, Stache-Crain B;  
 XX WPI; 2001-530177/58.

XX WPI; 2001-530177/58.

PT New polynucleotides and polypeptides, useful for diagnosis and  
 PT treatment of breast, lung and colon cancer -  
 XX

PS Claim 1; Page 896; 1193pp; English.

XX The invention relates to new polynucleotides and polypeptides, useful for  
 CC diagnosis and treatment of breast, lung and colon cancer. The sequences  
 CC can be used in detecting differentially expressed genes correlated with a  
 CC cancerous state of a mammalian cell, comprising detecting at least one  
 CC differentially expressed gene product in a test sample derived from a  
 CC cell suspected of being cancerous. They can also be used to inhibit  
 CC tumour growth by modulating expression of a gene product. AAS36943-  
 CC AAS3338 represent novel human diagnostic and therapeutic coding  
 CC sequences of the invention.

XX Sequence 408 BP, 57 A, 110 C, 140 G, 101 T, 0 other;

Query Match 70.0%; Score 14; DB 22; Length 408;

Best Local Similarity 100.0%; Pred. No. 1.3e+02;

Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 OY 7 GAGGACCAAGTCCC 20  
 DB 91 GAGGACCAAGTCCC 78

RESULT 31  
 AAX55494/c  
 ID AAX55494 standard; DNA; 409 BP.

XX AAX55494;

XX 27-JUL-1999 (first entry)

XX Rice Crm1 homologue DNA.

XX Pad1; Crm1; Jab1; Ap-1 transcription factor activity; regulator; plant;  
 KW maize; soybean; wheat; rice; yeast; human; isolation; transgenic; ss.  
 XX

XX Oryza sativa.

XX WO9924574-A2.

XX 20-MAY-1999.

XX 04-NOV-1998; 98MO-US23487.

XX 07-NOV-1997; 97US-0064914.

XX (DUPO ) DU PONT DE NEMOURS & CO E. I.

XX Allen SM, Anderson SL, Hitz WD, Kinney AJ, Miao G;  
 PI Morgante M, Odell JT, Sakai H;  
 XX WPI; 1999-327405/27.

XX P-FSDB; AAY08447.

PT Plant homologues of yeast Pad1, Crm1 and human Jab1 and related  
 PT polynucleotides

XX Claim 7; Page 42; 57pp; English.

XX This invention describes novel plant Pad1, Crm1 or Jab1 proteins which  
 CC are capable of Ap-1 transcription factor regulation. The proteins are  
 CC thought to interact with transcription factors altering gene expression.  
 CC The nucleic acid sequences of the invention may be used to isolate cDNAs  
 CC and genes encoding homologous proteins from the same or other plant  
 CC species. Synthetic peptides of the proteins may be synthesized to  
 CC generate antibodies that are useful for screening expression libraries.  
 CC Transgenic plants may be produced using the nucleic acid sequences to  
 CC alter the levels of Pad1, Crm1 and Jab1 present in the plants. Altering  
 CC the levels of these proteins would alter the level of Ap-1 transcription  
 CC factor activity in the plants.

SQ Sequence 409 BP, 119 A, 90 C, 82 G, 118 T, 0 other;

Query Match 70.0%; Score 14; DB 20; Length 409;

Best Local Similarity 100.0%; Pred. No. 1.3e+02;  
 Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 2 CTTAGGAGGACCA 15  
 DB 296 CTTAGGAGGACCA 283

RESULT 32  
 AAS26323  
 ID AAS26323 standard; cDNA; 442 BP.

XX AAS26323;

XX 07-NOV-2001 (first entry)

XX Human cDNA encoding a novel secreted protein, Seq ID 502.  
DE  
XX  
KW Human; immunosuppressive; antiarthritic; ss; antirheumatic;  
KW cytoskeletal; cardiant; vasotropic; cerebroprotective; nocitropic;  
KW neuroprotective; antibacterial; virucide; fungicide; ophthalmological;  
KW vulnary; secreted protein; rheumatoid arthritis;  
KW hyperproliferative disorder; cardiovascular disorder; cardiac arrest;  
KW cerebrovascular disorder; cerebral ischemia; angiogenesis;  
KW nervous system disorder; Alzheimer's disease; infection; ocular disorder;  
KW corneal infection; wound healing; epithelial cell proliferation;  
KW skin ageing; food additive; preservative; antiproliferative.  
OS Homo sapiens.  
XX  
PN W0200155322-A2.  
XX  
XX  
PD 02-AUG-2001.  
XX  
PF 17-JAN-2001; 2001WO-US01341.  
XX  
XX 31-JAN-2000; 2000US-0179065.  
PR 04-FEB-2000; 2000US-0180628.  
PR 24-FEB-2000; 2000US-0184664.  
PR 02-MAR-2000; 2000US-0186350.  
PR 16-MAR-2000; 2000US-0189874.  
PR 17-MAR-2000; 2000US-0190076.  
PR 18-APR-2000; 2000US-0198123.  
PR 19-MAY-2000; 2000US-0205515.  
PR 07-JUN-2000; 2000US-0209467.  
PR 28-JUN-2000; 2000US-0214886.  
PR 30-JUN-2000; 2000US-0215135.  
PR 07-JUL-2000; 2000US-0216647.  
PR 11-JUL-2000; 2000US-0216880.  
PR 11-JUL-2000; 2000US-0217487.  
PR 11-JUL-2000; 2000US-0217496.  
PR 14-JUL-2000; 2000US-0218290.  
PR 26-JUL-2000; 2000US-0220963.  
PR 26-JUL-2000; 2000US-0220964.  
PR 14-AUG-2000; 2000US-0224518.  
PR 14-AUG-2000; 2000US-0224519.  
PR 14-AUG-2000; 2000US-0225213.  
PR 14-AUG-2000; 2000US-0225214.  
PR 14-AUG-2000; 2000US-0225266.  
PR 14-AUG-2000; 2000US-0225267.  
PR 14-AUG-2000; 2000US-0225268.  
PR 14-AUG-2000; 2000US-0225270.  
PR 14-AUG-2000; 2000US-0225447.  
PR 14-AUG-2000; 2000US-0225757.  
PR 14-AUG-2000; 2000US-0225758.  
PR 14-AUG-2000; 2000US-0225759.  
PR 18-AUG-2000; 2000US-0226279.  
PR 22-AUG-2000; 2000US-0226681.  
PR 22-AUG-2000; 2000US-0226868.  
PR 22-AUG-2000; 2000US-0227182.  
PR 23-AUG-2000; 2000US-0227009.  
PR 30-AUG-2000; 2000US-0228924.  
PR 01-SEP-2000; 2000US-0229287.  
PR 01-SEP-2000; 2000US-0229343.  
PR 01-SEP-2000; 2000US-0229344.  
PR 01-SEP-2000; 2000US-0229345.  
PR 05-SEP-2000; 2000US-0229509.  
PR 05-SEP-2000; 2000US-0229513.  
PR 06-SEP-2000; 2000US-0230437.  
PR 06-SEP-2000; 2000US-0230438.  
PR 08-SEP-2000; 2000US-0231242.  
PR 08-SEP-2000; 2000US-0231243.  
PR 08-SEP-2000; 2000US-0231244.  
PR 08-SEP-2000; 2000US-0231413.  
PR 08-SEP-2000; 2000US-0231414.  
PR 08-SEP-2000; 2000US-0232080.  
PR 08-SEP-2000; 2000US-0232081.  
PR 12-SEP-2000; 2000US-0231968.  
PR 14-SEP-2000; 2000US-0232397.  
PR 14-SEP-2000; 2000US-0232398.  
PR 14-SEP-2000; 2000US-0232399.  
PR 14-SEP-2000; 2000US-0232400.  
PR 14-SEP-2000; 2000US-0232401.  
PR 14-SEP-2000; 2000US-0233063.  
PR 14-SEP-2000; 2000US-0233064.  
PR 14-SEP-2000; 2000US-0233065.  
PR 21-SEP-2000; 2000US-0234223.  
PR 21-SEP-2000; 2000US-0234274.  
PR 25-SEP-2000; 2000US-0234397.  
PR 25-SEP-2000; 2000US-0234998.  
PR 26-SEP-2000; 2000US-0235484.  
PR 27-SEP-2000; 2000US-0235834.  
PR 27-SEP-2000; 2000US-0235836.  
PR 29-SEP-2000; 2000US-0236327.  
PR 29-SEP-2000; 2000US-0236367.  
PR 29-SEP-2000; 2000US-0236368.  
PR 29-SEP-2000; 2000US-0236369.  
PR 29-SEP-2000; 2000US-0236370.  
PR 02-OCT-2000; 2000US-0237037.  
PR 02-OCT-2000; 2000US-0237038.  
PR 02-OCT-2000; 2000US-0237039.  
PR 02-OCT-2000; 2000US-0237040.  
PR 13-OCT-2000; 2000US-0239935.  
PR 13-OCT-2000; 2000US-0239937.  
PR 20-OCT-2000; 2000US-0240960.  
PR 20-OCT-2000; 2000US-0241221.  
PR 20-OCT-2000; 2000US-0241785.  
PR 20-OCT-2000; 2000US-0241786.  
PR 20-OCT-2000; 2000US-0241787.  
PR 20-OCT-2000; 2000US-0241808.  
PR 20-OCT-2000; 2000US-0241809.  
PR 20-OCT-2000; 2000US-0241826.  
PR 01-NOV-2000; 2000US-0244617.  
PR 08-NOV-2000; 2000US-0246474.  
PR 08-NOV-2000; 2000US-0246475.  
PR 08-NOV-2000; 2000US-0246476.  
PR 08-NOV-2000; 2000US-0246477.  
PR 08-NOV-2000; 2000US-0246478.  
PR 08-NOV-2000; 2000US-0246523.  
PR 08-NOV-2000; 2000US-0246524.  
PR 08-NOV-2000; 2000US-0246525.  
PR 08-NOV-2000; 2000US-0246526.  
PR 08-NOV-2000; 2000US-0246527.  
PR 08-NOV-2000; 2000US-0246528.  
PR 08-NOV-2000; 2000US-0246532.  
PR 08-NOV-2000; 2000US-0246609.  
PR 08-NOV-2000; 2000US-0246610.  
PR 08-NOV-2000; 2000US-0246611.  
PR 08-NOV-2000; 2000US-0246613.  
PR 17-NOV-2000; 2000US-0249207.  
PR 17-NOV-2000; 2000US-0249208.  
PR 17-NOV-2000; 2000US-0249209.  
PR 17-NOV-2000; 2000US-0249210.  
PR 17-NOV-2000; 2000US-0249211.  
PR 17-NOV-2000; 2000US-0249212.  
PR 17-NOV-2000; 2000US-0249213.  
PR 17-NOV-2000; 2000US-0249214.  
PR 17-NOV-2000; 2000US-0249215.  
PR 17-NOV-2000; 2000US-0249264.  
PR 17-NOV-2000; 2000US-0249265.  
PR 17-NOV-2000; 2000US-0249297.  
PR 17-NOV-2000; 2000US-0249299.  
PR 17-NOV-2000; 2000US-0249300.  
PR 01-DEC-2000; 2000US-0250160.  
PR 01-DEC-2000; 2000US-0250391.

PR 05-DEC-2000; 2000US-0251030.  
PR 05-DEC-2000; 2000US-0251988.  
PR 05-DEC-2000; 2000US-0256719.  
PR 06-DEC-2000; 2000US-0251479.  
PR 08-DEC-2000; 2000US-0251856.  
PR 08-DEC-2000; 2000US-0251868.  
PR 08-DEC-2000; 2000US-0251869.  
PR 08-DEC-2000; 2000US-0251989.  
PR 08-DEC-2000; 2000US-0251990.  
PR 11-DEC-2000; 2000US-0254097.  
PR 05-JAN-2001; 2001US-0259678.  
XX  
XX (HUMA-) HUMAN GENOME SCI INC.  
XX  
XX PI Rosen CA, Barash SC, Ruben SM;  
XX WPI, 2001-488783/53.  
XX P-PSDB; ABU16336.  
XX  
XX New nucleic acid molecules encoding 461 human secreted proteins for  
PT diagnosing, preventing, treating or ameliorating medical conditions and  
PT used as food additives or preservatives -  
XX  
XX Claim 1, SEQ ID No 502; 980bp; English.  
XX  
XX The invention relates to isolated nucleic acid molecules and their  
CC encoded secreted proteins. The nucleic acids and proteins are used to  
CC prevent, treat or ameliorate a medical condition in e.g. humans, mice,  
CC rabbits, goats, horses, cats, dogs, chickens or sheep. They  
CC are also used in diagnosing a pathological condition or susceptibility  
CC to a pathological condition. Antibodies to the proteins can also  
CC be used in alleviating symptoms associated with the disorders and in  
CC diagnostic immunoassays e.g. radioimmunoassays or enzyme linked  
CC immunoassays (ELISA). Disorders which are diagnosed or treated  
CC include autoimmune diseases e.g. Rheumatoid arthritis,  
CC hyperproliferative disorders e.g. neoplasms of the breast or liver,  
CC cardiovascular disorders e.g. cardiac arrest, cerebrovascular disorders  
CC e.g. cerebral ischaemia, angiogenesis, nervous system disorders e.g.  
CC Alzheimer's disease, infections caused by bacteria, viruses and fungi  
CC and ocular disorders e.g. corneal infection, and many other  
CC disorders listed in the specification. The polypeptides can also  
CC be used to aid wound healing and epithelial cell proliferation, to  
CC prevent skin aging due to sunburn, to maintain organs before  
CC transplantation, for supporting cell culture of primary tissues, to  
CC regenerate tissues and in chemotaxis. The polypeptides can also be used  
CC as a food additive or preservative to increase or decrease storage  
CC capabilities, fat content, lipid, protein, carbohydrate, vitamins,  
CC minerals, cofactors and other nutritional components. The present  
CC sequence encodes a novel secreted protein of the invention.

Query March 70.0%; Score 14; DB 22; Length 442;  
Best Local Similarity 100.0%; Pred. NO. 1.3e+02;  
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 4 TAGGAGGACAACT 17  
141 TAGGAGGACAACT 154  
Db

RESULT 33  
ABX73664  
ID ABX73664 standard; DNA; 442 BP.  
XX  
XX AC ABX73664;  
XX  
XX DT 18-MAR-2003 (first entry)  
XX  
XX DE Human novel polynucleotide #492.  
XX  
XX KW Human; gene; ds; neural disorder; immune system disorder; renal disorder;  
KW muscular disorder; respiratory disease; reproductive disorder;  
KW gastrointestinal disorder; pulmonary disorder; cardiovascular disorder;  
KW hyperproliferative disorder; inflammatory disease; allergic reaction;

KW blood related disorder; cancer; immunosuppressive; antiinflammatory;  
KW cardiovascular; nephrotoxic; cytostatic; antiallergic; chromolytic;  
KW haemostatic; antiarteriosclerotic.  
XX  
XX OS Homo sapiens.  
XX  
XX PN US2002132753-A1.  
XX  
XX PD 19-SEP-2002.  
XX  
XX PF 17-JAN-2001; 2001US-0764864.  
XX  
XX 31-JAN-2000; 2000US-179065P.  
XX 04-FEB-2000; 2000US-180628P.  
XX 28-JUN-2000; 2000US-214866P.  
XX 07-JUL-2000; 2000US-216647P.  
XX 07-JUL-2000; 2000US-216880P.  
XX 11-JUL-2000; 2000US-217487P.  
XX 14-JUL-2000; 2000US-218290P.  
XX 26-JUL-2000; 2000US-220963P.  
XX 26-JUL-2000; 2000US-220964P.  
XX 14-AUG-2000; 2000US-224518P.  
XX 14-AUG-2000; 2000US-224519P.  
XX 14-AUG-2000; 2000US-225267P.  
XX 14-AUG-2000; 2000US-225268P.  
XX 14-AUG-2000; 2000US-225270P.  
XX 14-AUG-2000; 2000US-225447P.  
XX 14-AUG-2000; 2000US-225757P.  
XX 14-AUG-2000; 2000US-225758P.  
XX 22-AUG-2000; 2000US-226868P.  
XX 30-AUG-2000; 2000US-228924P.  
XX 01-SEP-2000; 2000US-229287P.  
XX 01-SEP-2000; 2000US-229343P.  
XX 01-SEP-2000; 2000US-229344P.  
XX 01-SEP-2000; 2000US-229345P.  
XX 05-SEP-2000; 2000US-229509P.  
XX 05-SEP-2000; 2000US-229513P.  
XX 08-SEP-2000; 2000US-231413P.  
XX 21-SEP-2000; 2000US-234223P.  
XX 21-SEP-2000; 2000US-234274P.  
XX 25-SEP-2000; 2000US-234979P.  
XX 27-SEP-2000; 2000US-235834P.  
XX 29-SEP-2000; 2000US-236327P.  
XX 29-SEP-2000; 2000US-236367P.  
XX 29-SEP-2000; 2000US-236368P.  
XX 29-SEP-2000; 2000US-236369P.  
XX 29-SEP-2000; 2000US-236370P.  
XX 02-OCT-2000; 2000US-236802P.  
XX 02-OCT-2000; 2000US-237037P.  
XX 02-OCT-2000; 2000US-237038P.  
XX 02-OCT-2000; 2000US-237039P.  
XX 02-OCT-2000; 2000US-237040P.  
XX 13-OCT-2000; 2000US-239935P.  
XX 20-OCT-2000; 2000US-240960P.  
XX 20-OCT-2000; 2000US-241785P.  
XX 20-OCT-2000; 2000US-241809P.  
XX 01-NOV-2000; 2000US-244617P.  
XX 17-NOV-2000; 2000US-249299P.  
XX 08-DEC-2000; 2000US-251866P.  
XX 08-DEC-2000; 2000US-251868P.  
XX 08-DEC-2000; 2000US-251869P.  
XX 08-DEC-2000; 2000US-251869P.  
XX (ROSE/) ROSEN C A.  
XX (RUBEN/) RUBEN S M.  
XX (BARASH/) BARASH S C.  
XX  
XX PI Rosen CA, Ruben SM, Barash SC;  
XX  
XX WPI, 2003-147444/14.  
XX P-PSDB; ABU55404.  
XX  
XX New polypeptides and nucleic acids, useful in gene therapy for

PT treating, inhibiting or preventing e.g. neural, immune system,  
PT muscular, respiratory, reproductive, gastrointestinal, pulmonary,  
PT cardiovascular or renal disorders -  
XX  
PS Claim 1; SEQ ID NO 502; 402bp; English.  
XX  
CC The invention relates to human novel polypeptides and their associated  
CC polynucleotides. The polypeptides and polynucleotides are useful in gene  
CC therapy for treating, inhibiting or preventing neural disorders, immune  
CC system disorders (e.g. systemic lupus erythematosus, rheumatoid  
CC arthritis and multiple sclerosis), muscular disorders, respiratory  
CC diseases (e.g. nasal vestibulitis, nasal polyps and sinusitis),  
CC reproductive disorders, gastrointestinal disorders, pulmonary disorders,  
CC cardiovascular disorders (e.g. congenital heart defects, Ebstein's  
CC anomaly and hypoplastic left heart syndrome), renal disorders (e.g. acute  
CC kidney failure and end-stage renal disease), hyperproliferative disorders  
CC (e.g. Hodgkin's disease and leukaemia), inflammatory diseases (e.g.  
CC septic shock, bursitis and appendicitis), allergic reactions and  
CC conditions (e.g. asthma), blood related disorders (e.g. thrombosis,  
CC atherosclerosis and myocardial infarction) and cancerous diseases.  
CC Sequences ABX73173-ABX74167 represent human novel polynucleotides of the  
CC invention.  
XX  
SQ Sequence 442 BP; 192 A; 70 C; 83 G; 94 T; 3 other;  
XX  
Query Match 70.0%; Score 14; DB 25; Length 442;  
Best Local Similarity 100.0%; Pred. No. 1.3e+02;  
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
OY 4 TAGAGGAAACAAGT 17  
DB 141 TAGAGGAAACAAGT 154  
XX  
RESULT 34  
ABV36454  
ID ABV36454 standard; cDNA; 467 BP.  
XX  
AC ABV36454;  
XX  
DT 16-SEP-2002 (first entry)  
XX  
DE Human prostate expression marker cDNA 36445.  
XX  
KW Human; prostate cancer; cytostatic; carcinogen; pharmacodynamic marker;  
KW pharmacogenomic marker; gene; ss.  
XX  
OS Homo sapiens.  
XX  
PN MO200160860-A2.  
XX  
PD 23-AUG-2001.  
XX  
PF 20-FEB-2001; 2001WO-US05171.  
XX  
PR 17-FEB-2000; 2000US-183319P.  
XX  
PR 16-MAR-2000; 2000US-189862P.  
XX  
PR 25-MAY-2000; 2000US-207454P.  
XX  
PR 09-JUN-2000; 2000US-211314P.  
XX  
PR 18-JUL-2000; 2000US-219007P.  
XX  
PR 13-DEC-2000; 2000US-255281P.  
XX  
PA (MILL-) MILLENNIUM PREDICTIVE MEDICINE INC.  
XX  
PI Schlegel R, Endege WO, Nonahan JE;  
XX  
DR WPI; 2001-662795/76.  
XX  
PT Novel isolated nucleic acid molecule associated with cancerous state of  
PT prostate cells and correlating with presence of prostate cancer, useful  
PT for detecting presence of prostate cancer, stage of prostate cancer -  
XX  
PS Claim 1; Page 7534; 11750bp; English.

XX  
CC The invention relates to an isolated nucleic acid molecule (I) comprising  
CC a nucleotide sequence given in Tables 1-9 (ABV00010-ABV62213) of the  
CC specification or its complement. (I) is useful for:  
CC (a) assessing whether a patient is afflicted with prostate cancer;  
CC (b) monitoring the progression of prostate cancer in a patient;  
CC (c) assessing the efficacy of a test compound to inhibit prostate  
CC cancer in a patient;  
CC (d) assessing the efficacy of a therapy for inhibiting prostate cancer  
CC in a patient;  
CC (e) selecting a composition for inhibiting prostate cancer in a patient;  
CC (f) assessing the prostate cell carcinogenic potential of a compound;  
CC (g) determining whether prostate cancer has metastasized in a patient;  
CC (h) assessing the aggressiveness or indolence of prostate cancer in a  
CC patient;  
CC (I) is also useful as a pharmacodynamic or pharmacogenomic marker.  
XX  
SQ Sequence 467 BP; 148 A; 84 C; 117 G; 118 T; 0 other;  
XX  
Query Match 70.0%; Score 14; DB 23; Length 467;  
Best Local Similarity 100.0%; Pred. No. 1.3e+02;  
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
OY 5 AGGAGGAACAAGTC 18  
DB 181 AGGAGGAACAAGTC 194  
XX  
RESULT 35  
ABA43976/C  
ID ABA43976 standard; DNA; 477 BP.  
XX  
AC ABA43976;  
XX  
DT 01-FEB-2002 (first entry)  
XX  
DE Human breast cell single exon nucleic acid probe #2671.  
XX  
KW Human; microarray; single exon probe; gene expression; breast;  
KW disease; cancer; ss.  
XX  
OS Homo sapiens.  
XX  
PN MO200157271-A2.  
XX  
PD 09-AUG-2001.  
XX  
PF 30-JAN-2001; 2001WO-US00662.  
XX  
PR 04-FEB-2000; 2000US-0180312.  
XX  
PR 26-MAY-2000; 2000US-0207456.  
XX  
PR 30-JUN-2000; 2000US-0608408.  
XX  
PR 03-AUG-2000; 2000US-0632366.  
XX  
PR 21-SEP-2000; 2000US-0234687.  
XX  
PR 27-SEP-2000; 2000US-0236359.  
XX  
PR 04-OCT-2000; 2000GB-0024263.  
XX  
PA (MOLE-) MOLECULAR DYNAMICS INC.  
XX  
PI Penn SG, Hanzel DK, Chen W, Rank DR;  
XX  
DR WPI; 2001-496933/54.  
XX  
PT New spatially-addressable set of single exon nucleic acid probes,  
PT useful for measuring gene expression in sample derived from human  
PT breast, comprises number of single exon nucleic acid probes -  
XX  
PS Claim 1; SEQ ID NO 2671; 327bp + sequence listing; English.  
XX  
CC The invention relates to a spatially-addressable set of single exon  
CC nucleic acid probes for measuring gene expression in a sample derived  
CC from human breast and BT 474 cells. The method involves contacting  
CC the probes with a collection of detectably labelled nucleic acids

CC derived from mRNA of human breast, and then measuring the label  
 CC bound to each probe of the microarray. The probes are useful for  
 CC verifying the expression of regions of genomic DNA predicted to  
 CC encode proteins. They are useful for gene discovery, and for  
 CC determining predisposition and/or prognosing breast disease. Gene  
 CC expression analysis is useful for assessing the toxicity of chemical  
 CC agents on cells. The microarray of this invention presents a far greater  
 CC diversity of probes for measuring gene expression, with far less bias  
 CC than expressed sequence tag microarrays. The method is suitable for  
 CC rapid production of functional information from genomic sequence. The  
 CC present sequence is a single exon nucleic acid probe of the invention.  
 CC Note: The sequence data for this patent did not form part of the  
 CC printed specification, but was obtained in electronic format directly  
 CC from WIP0 at ftp.wipo.int/pub/published\_pct\_sequences.  
 XX

SQ Sequence 477 BP; 112 A; 93 C; 127 G; 145 T; 0 other;

Query Match 70.0%; Score 14; DB 22; Length 477;  
 Best Local Similarity 100.0%; Pred. No. 1.3e+02;

Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 7 GAGGACACATCCC 20  
 |||||

Db 314 GAGGACACATCCC 301

Search completed: August 14, 2003, 21:41:21  
 Job time : 130 secs

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GenCore version 5.1.6  
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: August 14, 2003, 21:41:37 ; Search time 1126.8 Seconds  
(without alignments)  
388.250 Million cell updates/sec

Title: US-10-074-620-1

Perfect score: 18  
Sequence: 1 ggcgcgtcaccctgta 18

Scoring table: OLIGO\_NUC  
Gapop 60.0 , Gapext 60.0

Searched: 22781392 seqs, 12152238056 residues

Word size : 0

Total number of hits satisfying chosen parameters: 45562784

Minimum DB seq length: 0  
Maximum DB seq length: 200000000

Post-processing: Listing first 120 summaries

Database :

EST:  
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2: em\_esthum:\*  
3: em\_estin:\*  
4: em\_estmu:\*  
5: em\_estov:\*  
6: em\_estpl:\*  
7: em\_estro:\*  
8: em\_hic:\*  
9: gb\_est1:\*  
10: gb\_est2:\*  
11: gb\_hic:\*  
12: gb\_est3:\*  
13: gb\_est4:\*  
14: gb\_est5:\*  
15: em\_estfun:\*  
16: em\_estcom:\*  
17: em\_gss\_hum:\*  
18: em\_gss\_inv:\*  
19: em\_gss\_pin:\*  
20: em\_gss\_vrt:\*  
21: em\_gss\_fun:\*  
22: em\_gss\_mam:\*  
23: em\_gss\_mus:\*  
24: em\_gss\_pro:\*  
25: em\_gss\_rtd:\*  
26: em\_gss\_phg:\*  
27: em\_gss\_vrt1:\*  
28: gb\_gss1:\*  
29: gb\_gss2:\*

Pred. No. is the number of results predicted by chance to have a  
score greater than or equal to the score of the result being printed,  
and is derived by analysis of the total score distribution.

# SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	17	94.4	927	10	BF536318
2	17	94.4	1080	29	CNS022AD
3	16	88.9	124	12	BI014294
4	16	88.9	170	9	AA294899

C 5	16	88.9	183	10	BF170207	BF170207 PCI0425 M
C 6	16	88.9	199	9	AM405433	AM405433 UI-HF-BLO
C 7	16	88.9	221	10	BF354113	BF354113 PM4-HT072
C 8	16	88.9	273	10	BE969911	BE969911 601679494
C 9	16	88.9	295	9	AA339056	AA339056 EST44112
C 10	16	88.9	300	9	AU099544	AU099544 AU099544
C 11	16	88.9	301	14	BF516504	BF516504 UI-H-BW1
C 12	16	88.9	317	10	CB267362	CB267362 1006268 H
C 13	16	88.9	319	13	BQ369040	BQ369040 PM3-GN051
C 14	16	88.9	319	13	BQ369218	BQ369218 PM3-GN051
C 15	16	88.9	335	10	BF850266	BF850266 CM3-EN007
C 16	16	88.9	377	14	CB130696	CB130696 K-EST0180
C 17	16	88.9	394	14	H69706	H69706 Y-93a08 .s1
C 18	16	88.9	395	12	BM789452	BM789452 K-EST0069
C 19	16	88.9	404	10	BF668175	BF668175 602122957
C 20	16	88.9	413	14	CB270357	CB270357 1009264 H
C 21	16	88.9	428	9	AA135078	AA135078 ZO26406 .r
C 22	16	88.9	430	9	AW732816	AW732816 DB14110 .y
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C 24	16	88.9	431	14	CB270361	CB270361 1009268 H
C 25	16	88.9	433	14	CB145604	CB145604 K-EST0200
C 26	16	88.9	434	14	CB129626	CB129626 K-EST0179
C 27	16	88.9	436	14	CB142213	CB142213 K-EST0195
C 28	16	88.9	438	10	BF858511	BF858511 RCL1-F1019
C 29	16	88.9	447	10	BG104409	BG104409 602311027
C 30	16	88.9	449	10	BF130453	BF130453 601818761
C 31	16	88.9	452	10	BE909128	BE909128 601501730
C 32	16	88.9	452	13	BK475327	BK475327 DKFZP686I
C 33	16	88.9	453	9	AA96839	AA96839 ae33b02 .r
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C 36	16	88.9	459	10	BF857906	BF857906 RCL1-F1019
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C 41	16	88.9	484	9	AW246279	AW246279 2821886 .5
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C 45	16	88.9	490	13	BK471656	BK471656 DKFZP686L
C 46	16	88.9	494	10	BF857889	BF857889 RCL1-F1019
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C 48	16	88.9	501	13	BK280514	BK280514 BX280514
C 49	16	88.9	509	10	BF971748	BF971748 602240020
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C 51	16	88.9	512	10	BF699171	BF699171 602126810
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C 53	16	88.9	520	9	AM328258	AM328258 dB01C08 .x
C 54	16	88.9	525	10	BE390761	BE390761 601287253
C 55	16	88.9	533	10	BE799191	BE799191 601592752
C 56	16	88.9	540	9	AA307552	AA307552 EST178632
C 57	16	88.9	542	14	CB144397	CB144397 K-EST0198
C 58	16	88.9	542	14	CB269294	CB269294 1008201 H
C 59	16	88.9	549	10	BG495444	BG495444 602539755
C 60	16	88.9	550	14	CB128792	CB128792 K-EST0178
C 61	16	88.9	551	13	BK475673	BK475673 DKFZP686H
C 62	16	88.9	551	14	CB266268	CB266268 1005173 H
C 63	16	88.9	557	10	BE389991	BE389991 601285456
C 64	16	88.9	561	10	BE266170	BE266170 601191319
C 65	16	88.9	570	10	BG687435	BG687435 602639402
C 66	16	88.9	572	9	AA203223	AA203223 x56602 .r
C 67	16	88.9	581	10	BG528181	BG528181 602557578
C 68	16	88.9	581	12	B1915162	B1915162 603177375
C 69	16	88.9	584	10	BE279400	BE279400 601157972
C 70	16	88.9	586	9	AM675194	AM675194 DB41a06 .y
C 71	16	88.9	588	10	BF701069	BF701069 602128125
C 72	16	88.9	592	12	BM048988	BM048988 603621859
C 73	16	88.9	596	10	BE733288	BE733288 601569464
C 74	16	88.9	602	10	BE298839	BE298839 601119850
C 75	16	88.9	603	10	BE266714	BE266714 601190278
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c 84 16 88.9 629 10 BG529643 BG529643 602558076
c 85 16 88.9 632 10 BB664539 BB664539 602704066
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c 87 16 88.9 639 10 BE741309 BE741309 601594146
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c 93 16 88.9 647 10 BF691465 BF691465 602247634
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c 96 16 88.9 651 10 BE958171 BE958171 601644629
c 97 16 88.9 652 14 Z99383 Z99383 HS299383 DK
c 98 16 88.9 653 10 BF106351 BF106351 601823571
c 99 16 88.9 655 10 BE548866 BE548866 601071888
c 100 16 88.9 656 10 BG284498 BG284498 602408640
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c 109 16 88.9 678 12 BG777022 BG777022 602664237
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c 114 16 88.9 687 10 BG496927 BG496927 602541345
c 115 16 88.9 692 10 BF667533 BF667533 602120890
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c 117 16 88.9 694 10 BF125491 BF125491 601763433
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c 119 16 88.9 703 12 B1258980 B1258980 602971630
c 120 16 88.9 705 10 BE277837 BE277837 601120048

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## ALIGNMENTS

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RESULT 1
BFS36318 927 bp mRNA linear EST 11-DEC-2000
LOCUS 602051968F1 NCI_CGAP_SG2 Mus musculus cDNA clone IMAGE:4191040 5',
DEFINITION mRNA sequence.
ACCESSION BFS36318 GI:11623686
VERSION BFS36318
KEYWORDS EST.
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus
REFERENCE NIH-MGC http://mgc.nci.nih.gov/
AUTHORS National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL Unpublished
COMMENT Contact: Robert Strausberg, Ph.D.
Email: CGAPs@mai.nih.gov
Tissue Procurement: Jeffrey E. Green, M.D.
CDNA Library Preparation: Life Technologies, Inc.
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LNL at:
http://image.llnl.gov

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FEATURES
source Plate: L1AM9519 row: h column: 17
High quality sequence scop: 686.
Location/Qualifiers
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/db_xref="taxon:10090"
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/lab_host="DH10B (T1 phage-resistant)"
/clone_lib="NCI_CGAP_SG2"
/note="Organ: salivary gland; Vector: PCMV-SPORE6; Site 1:
NotI; Site 2: SalI; Cloned unidirectionally. Primer: Oligo
dT. Average insert size 1.3 kb. Constructed by Life
Technologies. Note: this is a NCI CGAP Library."
BASE COUNT 221 a 224 c 266 g 214 t 2 others
ORIGIN
Query Match 94.4%; Score 17; DB 10; Length 927;
Best Local Similarity 100.0%; Pred. No. 22;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 2 GCTGTCACCTGTTA 18
Db 818 GCTGTCACCTGTTA 834
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CNS02AHD 1080 bp DNA linear GSS 01-SEP-2000
LOCUS Tetraodon nigroviridis genome survey sequence T7 end of clone
DEFINITION 25101 of library G from Tetraodon nigroviridis, genomic survey
sequence.
ACCESSION AL188554 GI:7826658
VERSION AL188554.1
KEYWORDS GSS; genome survey sequence.
SOURCE Tetraodon nigroviridis
ORGANISM Tetraodon nigroviridis
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Actinopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei;
Acanthomorpha; Acanthopterygii; Perciformes; Tetraodontiformes;
Tetraodontidae; Tetraodontidae; Tetraodon.
REFERENCE 1 Roest Crolius,H., Jallion,O., Dasilva,C., Bouneau,L., Fisher,C.,
Bernot,A., Fizames,C., Wincker,P., Brottier,P., Quetier,F.,
Saurin,M. and Weissenbach,U.
Estimate of human gene number provided by genome-wide analysis
using Tetraodon nigroviridis DNA sequence
Nat. Genet. 25 (2), 235-238 (2000)
JOURNAL 20296633
MEDLINE 10835645
REFERENCE 2
AUTHORS Roest Crolius,H., Jallion,O., Dasilva,C., Ozouf-Costaz,C.,
Fizames,C., Fischer,C., Bouneau,L., Billaule,A., Quetier,F.,
Saurin,M., Bernot,A. and Weissenbach,U.
Characterization and repeat analysis of the compact genome of the
freshwater pufferfish Tetraodon nigroviridis
Genome Res. 10 (7), 939-949 (2000)
JOURNAL 20359837
MEDLINE 10899143
REFERENCE 3 (bases 1 to 1080)
AUTHORS Genoscope.
TITLE Direct Submission
JOURNAL Submitted (12-APR-2000) Genoscope - Centre National de Sequencage :
BP 191 91006 EVRY cedex - FRANCE (E-mail : sequef@genoscope.cns.fr
Web : www.genoscope.cns.fr)
COMMENT This sequence is a single read and was generated as part of a large
scale clone-end sequencing project of the Tetraodon nigroviridis
genome. For more information, please take a look at
http://www.genoscope.cns.fr/Tetraodon.
FEATURES
source Location/Qualifiers
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Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Db 324 GCTGTGTCACCTGTTA 340

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LOCUS MR4-ET0142-310101-001-h11 ET0142 Homo sapiens cDNA, mRNA sequence.  
DEFINITION BI014294  
VERSION BI014294.1 GI:14418365  
KEYWORDS EST.  
SOURCE Homo sapiens (human)  
ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE 1 (bases 1 to 124)  
Dias Neto,E., Garcia Correa,R., Verjovski-Almeida,S., Brites,M.R.,  
Nagai,M.A., da Silva,W. Jr., Zago,M.A., Bordin,S., Costa,F.F.,  
Goldman,G.H., Carvalho,A.F., Matsukuma,A., Bala,G.S., Simpson,D.H.,  
Brunstein,A., deOliveira,P.S., Bucher,P., Jongeneel,C.V., O'Hare,  
M.J., Soares,F., Brentani,R.K., Reis,L.F., de Souza,S.J. and  
Simpson,A.U.

TITLE Shotgun sequencing of the human transcriptome with ORF expressed  
sequence tags  
JOURNAL Proc. Natl. Acad. Sci. U.S.A. 97 (7), 3491-3496 (2000)  
MEDLINE 2002663  
PUBMED 10737800  
COMMENT Contact: Simpson A.J.G.  
Laboratory of Cancer Genetics  
Ludwig Institute for Cancer Research  
Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP,  
Brazil  
Tel: +55-11-2704922  
Fax: +55-11-2707001  
Email: asimpson@ludwig.org.br  
This sequence was derived from the FAPESP/LICR Human Cancer Genome  
Project. This entry can be seen in the following URL.  
(http://www.ludwig.org.br/scripts/gethtml2.pl?tl=MR4&ct=MR4-ET0142-  
310101-001-h11&ct3=2001-01-31&ct4=1)  
Seq primer: puc 18 forward  
High quality sequence start: 16  
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Location/Qualifiers

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Site\_2: Sma1; A mini-library was made by cloning products  
derived from ORESTES PCR (U.S. Letters Patent application  
No. 196,716 - Ludwig Institute for Cancer Research)  
profiles into the pUC 18 vector. Reverse transcription of  
tissue mRNA and cDNA amplification were performed under  
low stringency conditions."

BASE COUNT 26 a 51 c 19 g 28 t  
ORIGIN

Query Match 88.9%; Score 16; DB 12; Length 124;

Best Local Similarity 100.0%; Pred. No. 47;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Db 90 GGCTGTGTCACCTGT 75

RESULT 4  
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LOCUS EST100142 Pancreas tumor I Homo sapiens cDNA 5' end, mRNA sequence.  
DEFINITION AA294899  
VERSION AA294899.1 GI:1947334  
KEYWORDS EST.  
SOURCE Homo sapiens (human)  
ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE 1 (bases 1 to 170)  
Adams,M.D., Kerlavage,A.R., Fleischmann,R.D., Fuldner,R.A., Bult  
C.J., Lee,N.H., Kirness,E.F., Weinstock,K.G., Gocayne,J.D., White  
O., Sutton,G., Blake,J.A., Brandon,R.C., Man-Wei,C., Clayton,R.A.,  
Cline,T.R., Cotton,M.D., Earle-Hughes,J., Fene,L.D., Fitzgerald,  
L.M., Fitzhugh,W.M., Fritchman,J.L., Geoghagen,N.S., Glodok,A.,  
Gnehm,C.L., Hanna,M.C., Hedblom,E., Hinkle,P.S., Jr., Kelley,J.M.,  
Kelley,J.C., Liu,L.-I., Marmaros,S.M., Merrick,J.M.,  
Moreno-Palantques,R.F., McDonald,L.A., Nguyen,D.T., Pelligrino,S.M.,  
Phillips,C.A., Ryder,S.E., Scott,U.L., Sauder,D.M., Shirley,R.,  
Small,K.V., Spriggs,T.A., Utterback,T.R., Weidman,J.F., Li,Y.,  
Bednarek,D.P., Cao,L., Cepeda,M.A., Coleman,T.A., Collins,E.J.,  
Dimke,D., Feng,D.-F., Ferris,A., Fischer,C., Hastings,G.A., He,W.W.,  
Hu,J.S., Greene,J.M., Gruber,J., Hudson,P., Kim,A.K., Kozak,D.L.,  
Kunesh,C., Hungjun,J., Li,H., Meisner,P.S., Olsen,H., Raymond,L.,  
Wei,Y.F., Wang,J., Xu,C., Yu,G.L., Ruben,S.M., Dillon,P.J., Fannon  
M.R., Rosen,C.A., Haseltine,W.A., Fields,C., Fraser,C.M. and  
Venter,J.C.

TITLE Initial assessment of human gene diversity and expression patterns  
based upon 83 million nucleotides of cDNA sequence  
JOURNAL Nature 377 (6547 Suppl), 3-174 (1995)  
MEDLINE 96026280  
PUBMED 7566098  
COMMENT Other ESTs: TH0172745  
Contact: Kerlavage, AR  
Bioinformatics  
The Institute for Genomic Research  
9712 Medical Center Drive, Rockville, MD 20850 USA  
Tel: 3018699056  
Fax: 3018699423  
Email: arkerlav@tigr.org  
For clone availability, additional sequence and expression  
information related to this EST, please check the TIGR Human Gene  
Index (http://www.tigr.org/cdb/hgi/hgi.html)  
Seq primer: M13 Reverse.

FEATURES  
source 1..170  
Location/Qualifiers

/organism="Homo sapiens"  
/mol\_type="mRNA"  
/db\_xref="ATCC (inhost):190475"  
/db\_xref="taxon:9606"  
/dev\_stage="adult"  
/clone\_1lb="Pancreas tumor I"  
/note="Organ: pancreas; Vector: pBluescript SK-; Site\_1:  
EcoRI, Site\_2: XhoI"

BASE COUNT 39 a 48 c 51 g 28 t 4 others  
ORIGIN

Query Match 88.9%; Score 16; DB 9; Length 170;  
Best Local Similarity 100.0%; Pred. No. 51;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGCTGTGTCACCTGT 16  
|||||

Db 135 GGCTGCTGCACCTGT 120

## RESULT 5

BF170207/c 183 bp mRNA linear EST 23-MAR-2001

LOCUS PC10425 Myeloma (PCL) cDNA library Homo sapiens CDNA, mRNA

## DEFINITION

sequence.

## ACCESSION

BF170207

## VERSION

BF170207.1 GI:13436309

## KEYWORDS

EST.

## SOURCE

ORGANISM

Homo sapiens (human)

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

1 (bases 1 to 183)

Woodgett,J.R. and Stewart,A.K.

'2.H., Nadeem,V., Cukerman,E., Francisco-Pabalan,O., Liew,C.C.,

Woodgett,J.R. and Stewart,A.K.

A molecular compendium of genes expressed in multiple myeloma

Blood 100 (6), 2175-2186 (2002)

22188429

12200383

COMMENT

Contact: A. Keith Stewart, M.D.

Oncology Research

University Health Network

610 University Ave., 5-126, Toronto, Ontario, M5G 2M9, Canada

Tel: (416) 946-6539

Fax: (416) 946-6546

Email: K.stewart@utoronto.ca

PCR Primers

FORWARD: 5'-GCCAAGCTCGAATTACCTCACTAAGG-3'

BACKWARD: 5'-CCAGTATGATGATGACTGACTGATGAGGCG-3'

Seq primer: 5'-GAATTAACCTCACTAAGG-3'.

Location/Qualifiers

1..183

/organism="Homo sapiens"

/mol\_type="mRNA"

/db\_xref="taxon:9606"

/sex="male"

/tissue\_type="Blood"

/cell\_type="myeloma"

/dev\_stage="Plasma cell leukemia"

/clone\_id="Myeloma (PCL) CDNA library"

/note="Vector: Lambda Zap Express; Site 1: EcoRI; Site 2:

XhoI; mRNA was purified from plasma cell leukemia

patient's peripheral blood containing >95% myeloma. An

oligo d(T)18 primer containing XhoI restriction site was

used to prime first strand synthesis using M-MuV reverse

transcriptase. To protect the cDNAs from XhoI digestion in

subsequent cloning step, the nucleotide analogue

5-methyl-dCTP was added to the nucleotide mixture and

la-32p(dATP) was added to monitor the quantity and quality

of first strand synthesis. After second-strand synthesis

and blunting of cDNA termini, EcoRI adapters were ligated

, followed by kinase treatment and digestion with XhoI.

The cDNAs were then size-fractionated using Sephacryl

S-500 column and then ligated into EcoRI and XhoI digested

Lambda Zap Express vector. The ligation product was

packaged using Gigapack II packaging extract. The library

had primary titre of approx. 1x10<sup>6</sup>. Clones from the

primary library were randomly selected for single pass

sequencing."

BASE COUNT 41 a 49 c 56 g 37 t

## ORIGIN

Query Match 88.9%; Score 16; DB 10; Length 183;

Best Local Similarity 100.0%; Pred. No. 52;

Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGCTGCTGCACCTGT 16

Db 143 GGCTGCTGCACCTGT 128

## RESULT 6

AM405433/c

LOCUS

DEFINITION UI-HF-BL0-adc-h-06-0-UI.r1 NIH\_MGC\_37 Homo sapiens CDNA clone

sequence.

IMAGE:3061115 5', mRNA sequence.

AM405433

VERSION

AM405433.1 GI:6924490

KEYWORDS

EST.

SOURCE

ORGANISM

Homo sapiens (human)

Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

1 (bases 1 to 199)

NIH-MGC http://mgs.nci.nih.gov/

National Institutes of Health, Mammalian Gene Collection (MGC)

Unpublished

CONTACT: Robert Strausberg, Ph.D.

Email: cgabs-r@mail.nih.gov

Eco RI site shown at the beginning of the sequence.

Tissue Procurement: Louis M. Staudt, M.D., Ph.D.

CDNA Library Preparation: M.B. Soares Lab

CDNA Library Arrayed by: M.B. Soares Lab

DNA Sequencing by: M.B. Soares Lab

Clone distribution: MGC clone distribution information can be

found through the I.M.A.G.E. Consortium/LLNL at:

www-bio.llnl.gov/bdrip/image/image.html

Seq primer: M13 Forward.

Location/Qualifiers

1..199

/organism="Homo sapiens"

/mol\_type="mRNA"

/db\_xref="taxon:9606"

/clone="IMAGE:3061115"

/tissue\_type="lymph"

/cell\_type="germinal center B cells"

/cell\_line="MGC85"

/lab\_host="DH10B (LT1)"

/clone\_id="NIH\_MGC\_37"

/note="Vector: pT73-Pac; Site 1: NotI; Site 2: Eco RI;

Constructed from size fractionated cytoplasmic mRNA

(1.5-2.5kb). Directionally cloned. Cells provided by Louis

M. Staudt, Ph.D. Library preparation by Maria de Fatima

Bonaldi, Ph.D. and M. Bento Soares, Ph.D."

BASE COUNT 60 a 57 c 53 g 29 t

## ORIGIN

Query Match 88.9%; Score 16; DB 9; Length 199;

Best Local Similarity 100.0%; Pred. No. 54;

Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGCTGCTGCACCTGT 16

Db 70 GGCTGCTGCACCTGT 55

## RESULT 7

BF354113/c

LOCUS

DEFINITION BF354113 221 bp mRNA linear EST 22-NOV-2000

PM4-HT0726-270500-002-c07 HT0726 Homo sapiens CDNA, mRNA sequence.

BF354113

VERSION

BF354113.1 GI:11313187

KEYWORDS

EST.

SOURCE

ORGANISM

Homo sapiens (human)

Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

1 (bases 1 to 221)

Dias Neto,E., Garcia Correa,R., Verjovski-Almeida,S., Briones,M.R.,

Nagai,M.A., da Silva,W. Jr., Zago,M.A., Bordin,S., Costa,F.F.,

Goldman,G.H., Carvalho,A.F., Matsukuma,A., Bala,G.S., Simpson,D.H.,

Brunstein,A., deOliveira,P.S., Bucher,P., Jongeneel,C.V., O'Hare

TITLE  
Simpson, A.J., Soares, F., Brentani, R.R., Reis, L.F., de Souza, S.J. and  
Simpson sequencing of the human transcriptome with ORF expressed  
sequence tags  
JOURNAL  
Proc. Natl. Acad. Sci. U.S.A. 97 (7), 3491-3496 (2000)  
MEDLINE  
200202663  
PUBMED  
10737800  
COMMENT  
Contact: Simpson A.J.G.

Laboratory of Cancer Genetics  
Ludwig Institute for Cancer Research  
Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP,  
Brazil  
Tel: +55-11-2704922  
Fax: +55-11-2707001  
Email: [astimmons@ludwig.org.br](mailto:astimmons@ludwig.org.br)  
This sequence was derived from the FAPSP/LICR Human Cancer Genome  
Project. This entry can be seen in the following URL  
(<http://www.ludwig.org.br/scripts/gethtml2.pl?c1=PM4&c2=PM4-HT0726>  
270500-002-c07&c3=2000-05-27&c4=1)  
Seq primer: puc 18 forward  
High quality sequence start: 28  
High quality sequence stop: 219.

FEATURES	SOURCE
location/Qualifiers	1. .221
/organism="Homo sapiens"	
/mol_type="mRNA"	
/db_xref="taxon:9606"	
/dev_stage="Adult"	
/clone_idb="HT0726"	
/note="Organ: head neck; Vector: puc18; Site 1: SmaI; Site 2: SmaI; A mini-library was made by cloning products derived from ORESTES PCR (U.S. Letters Patent application No. 196,716 - Ludwig Institute for Cancer Research) tissue mRNA and cDNA amplification were performed under low stringency conditions."	
BASE COUNT	39 a 94 c 31 g 57 t
ORIGIN	

Query Match	88.9%	Score 16;	DB 10;	Length 221;
Best Local Similarity	100.0%	Pred. No. 55;		
Matches	16;	Conservative	0;	Mismatches 0;
				Indels 0;
				Gaps 0;
QY	1	GGCTGGTGTCACTGT	16	
DB	99	GGCTGGTGTCACTGT	84	.

LOCUS	BE669911	273 bp	mRNA	linear	EST 04-OCT-2000
DEFINITION	601679494p1 NIH_MGC_78 Homo sapiens cDNA clone IMAGE:3949366 5', mRNA sequence.				
ACCESSION	BE669911				
VERSION	BE669911.1	GI:10582844			
KEYWORDS	EST.				
SOURCE	Homo sapiens (human)				
ORGANISM	Homo sapiens				
REFERENCE	Mullalyota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Eumetazoa; Eutheria; Primates; Carnivora; Homnidae; Homo.				
AUTHORS	1 (bases 1 to 273)				
TITLE	NIH-MGC <a href="http://mgc.nci.nih.gov/">http://mgc.nci.nih.gov/</a> .				
JOURNAL	National Institutes of Health, Mammalian Gene Collection (MGC)				
COMMENT	Unpublished				
	Contact: Robert Strausberg, Ph.D.				

REFERENCE	AUTHORS	TITLE	JOURNAL	COMMENT
1 (MGC 1 to 273)	NIH-MGC	<a href="http://mgc.nci.nih.gov/">http://mgc.nci.nih.gov/</a>	National Institutes of Health, Mammalian Gene Collection (MGC)	Unpublished
	Contact: Robert Strausberg, Ph.D.			
	Email: <a href="mailto:cgabbs@email.nih.gov">cgabbs@email.nih.gov</a>			
	Tissue Procurement: CLONETECH Laboratories, Inc.			
	cDNA Library Preparation: CLONETECH Laboratories, Inc.			
	cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNL)			
	DNA Sequencing by: Incyte Genomics, Inc.			
	Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LNL at:			
	<a href="http://image.llnl.gov">http://image.llnl.gov</a>			

```

plate: L1CM815 row: k column: 23
High quality sequence stop: 271.
Location/Qualifiers
FEATURES
    source      1..273

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/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:3949966"
/lab_host="DH10B (T1 phage-resistant)"
/clone_id="NH1_MGC_78"
/notes="Organ: pancreas; Vector: pDNR-LIB (Clontech);
Site_1: Sfil (ggcgccctcgcc); Site_2: Sfil (ggcattatggcc
); 5' and 3' adaptors were used in cloning as follows: 5'
adaptor sequence: 5'-CACGGCCATTATGGCC-3' and 3' adaptor
sequence: 5'-ATTCTAGAGCCGCGGCGGCACATG-dT(30)BN-3',
(where B = A, C, G or T). Average
insert size 1.2 kb (range 0.5-4.0 kb). 14/15 colonies
contained inserts by PCR. This library was enriched for
full-length clones and was constructed by Clontech
Laboratories (Palo Alto, CA)."

```

Query March	88.9%	Score 16	DB 10	Length 273
Best Local Similarity	100.0%	Pred. No. 58		
Matches 16	Conservative 0	Mismatches 0	Indels 0	Gaps 0
QY	1	GGCTGGGTCTCACCCTGT	16	
Db	151	GGCTGGGTCTCACCCTGT	136	

RESULT 9	AA339056/c	LOCUS	AA339056	295 bp	mRNA	linear	EST 21-APR-1997
DEFINITION	AA344112	Fetal brain I Homo sapiens	cDNA 5' end, mRNA sequence.				
ACCSSION	AA339056						
VERSION	AA339056.1	GI:1991304					
KEYWORDS	EST.						
SOURCE	Homo sapiens (human)						
ORGANISM	Homo sapiens						
	Eukaryota; Metazoa; Chordata; Cranialta; Vertebrata; Euteleostomi;						
	Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.						
REFERENCE	1 (bases 1 to 295)						
AUTHORS	Adams,M.D., Kerlavage,A.R., Fleischmann,R.D., Fuldner,R.A., Bult						

TITLE	Initial assessment of human gene diversity and expression patterns based upon 83 million nucleotides of cDNA sequence
JOURNAL	Nature 377 (6547 Suppl), 3-174 (1995)
MEDLINE	96026280
PUBMED	7566098
COMMENT	Other ESTs: THC172745

The Institute for Genomic Research  
9712 Medical Center Drive, Rockville, MD 20850 USA  
Tel: 3018699056  
Fax: 3018699423



Query Match 88.9%; Score 16; DB 10; Length 301;  
 Best Local Similarity 100.0%; Pred. No. 60;  
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GGCTGGTGCACCTGT 16  
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 Db 229 GGCTGGTGCACCTGT 244  
 |||||

RESULT 12  
 CB267362 317 bp mRNA linear EST 20-FEB-2003.  
 LOCUS CB267362/c  
 DEFINITION 1006268 Human Fat Cell 5'-Stretch Plus cDNA Library Homo sapiens  
 CDNA 5', mRNA sequence.  
 ACCESSION CB267362  
 VERSION CB267362.1 GI:28441947  
 KEYWORDS EST.  
 SOURCE Homo sapiens (human)  
 ORGANISM Homo sapiens  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.  
 1 (bases 1 to 317)  
 Yang, R.-Z., Shuldiner, A. and Gong, D.-W.  
 EST analysis of human adipose gene expression  
 JOURNAL Unpublished  
 CONTACT: Gong Da-Mei  
 Division of Endocrinology, Diabetes and Nutrition  
 University of Maryland  
 660 Redwood St., HH497, Baltimore, MD 21201, USA  
 Tel: 410 706 1672  
 Fax: 410 706 1622  
 Email: dgong@medicine.umaryland.edu  
 PCR Primers  
 FORWARD: CTCGGAGACGCCCATGTTGTTGTT  
 BACKWARD: AATAGACTACTATAGGCGCATTTGG  
 Seq primer: GTTGGTACCGGGAATTC.  
 Location/Qualifiers  
 1..317  
 /organism="Homo sapiens"  
 /mol\_type="mRNA"  
 /db\_xref="taxon:9606"  
 /sex="Male and Female"  
 /tissue\_type="adipose"  
 /clone\_id="Human Fat Cell 5'-Stretch Plus cDNA Library"  
 /note="Vector: lambda diriplex"

BASE COUNT 89 a 78 c 97 g 52 t 1 others

ORIGIN

Query Match 88.9%; Score 16; DB 14; Length 317;  
 Best Local Similarity 100.0%; Pred. No. 61;  
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GGCTGGTGCACCTGT 16  
 |||||  
 Db 136 GGCTGGTGCACCTGT 121  
 |||||

RESULT 13  
 BQ369040 319 bp mRNA linear EST 21-MAY-2002  
 LOCUS BQ369040/c  
 DEFINITION PM3-GN0516-060601-031-g03 GN0516 Homo sapiens cDNA, mRNA sequence.  
 ACCESSION BQ369040  
 VERSION BQ369040.1 GI:21044554  
 KEYWORDS EST.  
 SOURCE Homo sapiens (human)  
 ORGANISM Homo sapiens  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.  
 1 (bases 1 to 319)  
 Dias Neto, E., Garcia Correa, R., Verjowski-Almeida, S., Briones, M.R.,  
 Nagai, M.A., da Silva, W. Jr., Zago, M.A., Bordin, S., Costa, F.F.,  
 Goldman, G.H., Carvalho, A.F., Matsukuma, A., Bala, G.S., Simpson, D.H.,

TITLE  
 JOURNAL  
 MEDLINE  
 PUBMED  
 COMMENT

CONTACT: Simpson A.J.G.  
 Laboratory of Cancer Genetics  
 Ludwig Institute for Cancer Research  
 Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP,  
 Brazil  
 Tel: +55-11-2704922  
 Fax: +55-11-2707001  
 Email: asimpson@ludwig.org.br  
 This sequence was derived from the PAPESP/LICR Human Cancer Genome  
 Project. This entry can be seen in the following URL  
 (<http://www.ludwig.org.br/scripts/gethtml2.pl?l=PM3-GN0516-060601-031-g03&t3=2001-06-06&t4=1>)  
 Seq primer: puc 18 forward  
 High quality sequence start: 17  
 High quality sequence stop: 319.  
 Location/Qualifiers  
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 /mol\_type="mRNA"  
 /db\_xref="taxon:9606"  
 /dev\_stage="Adult"  
 /clone\_id="GN0516"  
 /note="Organ: placenta normal; Vector: puc18; Site 1: SmaI  
 products derived from ONESIES PCR (U.S. Letters Patent  
 application No. 196,716 - Ludwig Institute for Cancer  
 Research) profiles into the puc 18 vector. Reverse  
 transcription of tissue mRNA and cDNA amplification were  
 performed under low stringency conditions."

BASE COUNT 93 a 79 c 92 g 54 t 1 others

ORIGIN

Query Match 88.9%; Score 16; DB 13; Length 319;  
 Best Local Similarity 100.0%; Pred. No. 61;  
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GGCTGGTGCACCTGT 16  
 |||||  
 Db 97 GGCTGGTGCACCTGT 82  
 |||||

RESULT 14  
 BQ369218/c 319 bp mRNA linear EST 21-MAY-2002  
 LOCUS BQ369218  
 DEFINITION PM3-GN0516-020701-031-g03 GN0516 Homo sapiens cDNA, mRNA sequence.  
 ACCESSION BQ369218  
 VERSION BQ369218.1 GI:21044732  
 KEYWORDS EST.  
 SOURCE Homo sapiens (human)  
 ORGANISM Homo sapiens  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.  
 1 (bases 1 to 319)  
 Dias Neto, E., Garcia Correa, R., Verjowski-Almeida, S., Briones, M.R.,  
 Nagai, M.A., da Silva, W. Jr., Zago, M.A., Bordin, S., Costa, F.F.,  
 Goldman, G.H., Carvalho, A.F., Matsukuma, A., Bala, G.S., Simpson, D.H.,  
 Brunstein, A., de Oliveira, P.S., Bucher, P., Jongeneel, C.V., O'Hare,  
 M.J., Soares, F., Brentani, R.R., Reis, L.F., de Souza, S.J. and  
 Simpson, A.J.G.  
 Shotgun sequencing of the human transcriptome with ORF expressed  
 sequence tags  
 JOURNAL Proc. Natl. Acad. Sci. U.S.A. 97 (7), 3491-3496 (2000)  
 MEDLINE  
 PUBMED  
 COMMENT

CONTACT: Simpson A.J.G.

Laboratory of Cancer Genetics  
Ludwig Institute for Cancer Research  
Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP,  
Brazil  
Tel: +55-11-2704922  
Fax: +55-11-2707001  
Email: asimpson@ludwig.org.br  
This sequence was derived from the PAPSP/LICR Human Cancer Genome  
Project. This entry can be seen in the following URL  
(http://www.ludwig.org.br/scripts/gethtml2.pl?tl=PM3&t2=PM3-GN0516-  
020701-031-g03&t3=2001-07-02&t4=1)  
Seq primer: puc 18 forward  
High quality sequence start: 17  
High quality sequence stop: 319.  
Location/Qualifiers

## FEATURES

source

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1..319
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/dev_stage="Adult"
/clone_1lb="GN0516"
/note="Organ: placenta normal; Vector: puc18; Site_1: SmaI
; Site_2: SmaI; A mini-library was made by cloning
products derived from ORESTES PCR (U.S. Letters Patent
application No. 196,716 - Ludwig Institute for Cancer
Research) profiles into the puc 18 vector. Reverse
transcription of tissue mRNA and cDNA amplification were
performed under low stringency conditions."
```

## BASE COUNT

93 a 79 c 92 g 54 t 1 others

## ORIGIN

Query Match 88.9%; Score 16; DB 13; Length 319;  
Best Local Similarity 100.0%; Pred. No. 61;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
Qy 1 GGCTGGTGCACCTGT 16  
Db 97 GGCTGGTGCACCTGT 82

RESULT 15  
BF850266/c 335 bp mRNA linear EST 16-JAN-2001  
LOCUS CM3-EN0077-181100-489-d09 EN0077 Homo sapiens cDNA, mRNA sequence.  
DEFINITION BF850266  
ACCESSION BF850266  
VERSION BF850266.1 GI:12237428  
KEYWORDS EST.  
SOURCE Homo sapiens (human)  
ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.  
1 (bases 1 to 335)

## REFERENCE

AUTHORS

Dias, Neco, E., Garcia Correa, R., Verjowski-Almeida, S., Briones, M.R.,  
Nagai, M.A., da Silva, W. Jr., Zago, M.A., Bordin, S., Costa, F.F.,  
Goldman, G.H., Carvalho, A.F., Matsukuma, A., Bala, G.S., Simpson, D.H.,  
Brunstein, A., de Oliveira, P.S., Bucher, P., Jongeneel, C.V., O'Hare,  
M.J., Soares, F., Brentani, R.R., Reis, L.F., de Souza, S.J. and  
Simpson, A.J.

## TITLE

Shotgun sequencing of the human transcriptome with ORF expressed  
sequence tags

JOURNAL Proc. Natl. Acad. Sci. U.S.A. 97 (7), 3491-3496 (2000)

MEDLINE 20202663

PUBMED 10737800

## COMMENT

Contact: Simpson A.J.G.  
Laboratory of Cancer Genetics  
Ludwig Institute for Cancer Research  
Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP,  
Brazil  
Tel: +55-11-2704922  
Fax: +55-11-2707001  
Email: asimpson@ludwig.org.br  
This sequence was derived from the PAPSP/LICR Human Cancer Genome  
Project. This entry can be seen in the following URL

(http://www.ludwig.org.br/scripts/gethtml2.pl?tl=CM3&t2=CM3-EN0077-  
181100-489-d09&t3=2000-11-18&t4=1)  
Seq primer: puc 18 forward  
High quality sequence start: 52  
High quality sequence stop: 335.  
Location/Qualifiers

## FEATURES

source

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1..335
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/dev_stage="Adult"
/clone_1lb="EN0077"
/note="Organ: lung normal; Vector: puc18; Site_1: SmaI;
Site_2: SmaI; A mini-library was made by cloning products
derived from ORESTES PCR (U.S. Letters Patent application
No. 196,716 - Ludwig Institute for Cancer Research)
profiles into the puc 18 vector. Reverse transcription of
tissue mRNA and cDNA amplification were performed under
low stringency conditions."
```

## BASE COUNT

69 a 96 c 107 g 63 t

## ORIGIN

Query Match 88.9%; Score 16; DB 10; Length 335;  
Best Local Similarity 100.0%; Pred. No. 62;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
Qy 1 GGCTGGTGCACCTGT 16  
Db 335 GGCTGGTGCACCTGT 320

RESULT 16  
CB130696/c 377 bp mRNA linear EST 29-JAN-2003  
LOCUS K-EST0180649 L11SNJ354 Homo sapiens cDNA clone L11SNJ354-17-C06 5',  
DEFINITION mRNA sequence.  
ACCESSION CB130696  
VERSION CB130696.1 GI:28095037  
KEYWORDS EST.  
SOURCE Homo sapiens (human)  
ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.  
1 (bases 1 to 377)

## REFERENCE

AUTHORS

Oh, K.J., Cheong, J.E., Sohn, H.Y., Kim, J.M., Park, H.S., Kim, S. and  
Kim, Y.S.  
21C Frontier Korean EST Project 2001  
Unpublished  
Contact: Kim YS  
Genome Research Center  
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52 Eoen-dong Yuseong-gu, Daejeon 305-333, South Korea  
Tel: +82-42-860-4470  
Fax: +82-42-860-4409  
Email: yongsung@mail.kribb.re.kr  
Plate: 17 row: C column: 06  
High quality sequence stop: 377.  
Location/Qualifiers

## FEATURES

source

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1..377
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="L11SNJ354-17-C06"
/sex="M"
/tissue_type="Liver"
/cell_type="Polygonal"
/cell_line="SNJ-354"
/lab_host="Top10F"
/clone_1lb="L11SNJ354"
/note="Organ: Liver; Vector: pCNS-D2; Site_1: EcoRI;
Site_2: NotI; The poly (A) + RNA was dephosphorylated with
bacterial alkaline phosphatase (BAP) and then decapped
```





Best Local Similarity 100.0%; Pred. No. 65;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGCTGGTGCACCTGT 16  
|||||  
Db 111 GGCTGGTGCACCTGT 96

RESULT 19  
BF668175/c 404 bp mRNA linear EST 21-DEC-2000  
LOCUS 602122957F1 NIH\_MGC\_56 Homo sapiens cDNA clone IMAGE:4279958 5',  
DEFINITION mRNA sequence.

ACCESSION BF668175  
VERSION BF668175.1 GI:11942070  
KEYWORDS EST.  
SOURCE Homo sapiens (human)

ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE NIH-MGC http://mgi.nci.nih.gov/  
1 (bases 1 to 404)

AUTHORS National Institutes of Health, Mammalian Gene Collection (MGC)  
TITLE Unpublished  
JOURNAL Contact: Robert Strausberg, Ph.D.  
COMMENT Email: cgabs-remail.nih.gov  
Tissue Procurement: ATCC

CDNA Library Preparation: CLONTECH Laboratories, Inc.  
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNL)

DNA Sequencing by: Incyte Genomics, Inc.

Clone distribution: MGC clone distribution information can be  
found through the I.M.A.G.E. Consortium/LNL at:

http://image.llnl.gov

Plate: LHCMI06 row: a column: 15

High quality sequence stop: 404.

Location/Qualifiers

## FEATURES

source

1. .404  
/organism="Homo sapiens"

/mol\_type="mRNA"

/db\_xref="taxon:9606"

/clone="IMAGE:4279958"

/rissue\_type="primitive neuroectoderm"

/lab\_host="DH10B (T1 phage-resistant)"

/clone\_11b="NIH MGC 56"

/note="Organ: brain; Vector: pDNR-LIB (Clontech); Site\_1:

Site1 (ggcgcctggcc); Site\_2: Site1 (ggcgcctggcc);

Double-stranded cDNA was prepared from cell line RNA. 5'

and 3' adaptors were used in cloning as follows: 5'

adaptor sequence: 5'-CACGGCCATTATGGCC-3' and 3' adaptor

sequence: 5'-ATCTAGAGCGCGCGCGCGCATG-dT(30)BN-3';

(where B = A, C or G and N = A, C, G or T). Average

insert size 1.65 kb (range 0.9-4.0 kb). 15/15 colonies

contained inserts by PCR. This library was enriched for

full-length clones and was constructed by Clontech

Laboratories (Palo Alto, CA)."

BASE COUNT 108 a 103 c 125 g 67 t 1 others

## ORIGIN

Query Match

Best Local Similarity 100.0%; Pred. No. 65;

Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGCTGGTGCACCTGT 16  
|||||

Db 154 GGCTGGTGCACCTGT 139

RESULT 20  
CB270357 413 bp mRNA linear EST 20-FEB-2003  
LOCUS 1009264 Human Fat Cell 5'-Stretch Plus cDNA Library Homo sapiens  
DEFINITION cDNA 5', mRNA sequence.

ACCESSION CB270357

VERSION CB270357

KEYWORDS

SOURCE

ORGANISM

VERSION CB270357.1 GI:28444942  
EST.

## KEYWORDS

SOURCE

ORGANISM

Homo sapiens (human)

Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE

AUTHORS

Yang, R.-Z., Shuldiner, A. and Gong, D.-W.

1 (bases 1 to 413)

EST analysis of human adipose gene expression

Unpublished

JOURNAL

COMMENT

Contact: Gong Da-Wei

Division of Endocrinology, Diabetes and Nutrition

University of Maryland

660 Redwood St., HR497, Baltimore, MD 21201, USA

Tel: 410 706 1672

Fax: 410 706 1622

Email: dgong@medicine.umaryland.edu

PCR Primers

FORWARD: CTCGGAGAGCGCCATTGTGTGTG

BACKWARD: AATAGACTACTATAGGCGCAATTGG

Seq primer: GTTGTACCCGGGAATTC.

Location/Qualifiers

1. .413  
/organism="Homo sapiens"

/mol\_type="mRNA"

/db\_xref="taxon:9606"

/sex="Male and Female"

/rissue\_type="adipose"

/clone\_11b="Human Fat Cell 5'-Stretch Plus cDNA Library"

/note="Vector: lambdaTriplex"

BASE COUNT 114 a 104 c 119 g 71 t 5 others

## ORIGIN

Query Match

Best Local Similarity 100.0%; Pred. No. 65;

Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGCTGGTGCACCTGT 16  
|||||

Db 131 GGCTGGTGCACCTGT 116

RESULT 21  
AA135078 428 bp mRNA linear EST 06-AUG-1997  
LOCUS 2026d06.r1 StrataGene colon (#937204) Homo sapiens cDNA clone  
DEFINITION IMAGE:588011 5', mRNA sequence.

ACCESSION AA135078  
VERSION AA135078.1 GI:1696180  
KEYWORDS EST.

SOURCE Homo sapiens (human)

ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE 1 (bases 1 to 428)  
Hillier, L., Lennon, G., Becker, M., Bonaldo, M.F., Chapelli, B.,  
Chissoe, S., Dietrich, N., Dubuque, T., Pavello, A., Gish, W., Hawkins,  
'M., Hultman, M., Kucada, T., Lacy, M., Le, M., Le, N., Mardis, E., Moore,  
'B., Morris, M., Parsons, J., Prange, C., Rikkin, L., Rohlfing, T.,  
Schellenberg, K., Soares, M.B., Tan, F., Thierry-Mieg, J., Trevaaskis, B.,  
Underwood, K., Woldmann, P., Waterston, R., Wilson, R. and Marra, M.

Generation and analysis of 280,000 human expressed sequence tags  
Genome Res. 6 (9), 807-828 (1996)

TITLE JOURNAL  
MEDLINE 97044478  
PUBMED 8889549

COMMENT Contact: Wilson RK  
Washington University School of Medicine  
444 Forest Park Parkway, Box 8501, St. Louis, MO 63108  
Tel: 314 286 1800  
Fax: 314 286 1810  
Email: estewatson.wustl.edu  
This clone is available royalty-free through LNL; contact the  
IMAGE Consortium (info@image.llnl.gov) for further information.



Insert Length: 1117 Std Error: 0.00  
Seq primer: -28M13 rev2 from Amersham  
High quality sequence stop: 263.  
Location/Qualifiers

## FEATURES

SOURCE

1.428  
/organism="Homo sapiens"  
/mol\_type="mRNA"  
/db\_xref="GDB:4620313"  
/db\_xref="taxon:9606"  
/clone="IMAGE:588011"  
/issue\_type="tumor"  
/cell\_line="T84 carcinoma cell line"  
/lab\_host="SOLR cells (kanamycin resistant)"  
/clone\_lib="Stratagene colon (#37204)"  
/note="Organ: colon; Vector: pBluescript SK-; Site\_1: EcoRI; Site\_2: XhoI; Cloned unidirectionally. Primer: Oligo dt: T-84 colonic epithelial cell line. Average insert size: 1.0 kb; Uni-ZAP XR Vector; -5' adaptor sequence: 5' GAATTCGCGACGAG 3' -3' adaptor sequence: 5' CTCGAGTTTCTTTTCTTTT 3'."  
BASE COUNT 128 a 112 c 113 g 71 t 4 others  
ORIGIN

Query Match 88.9%; Score 16; DB 9; Length 428;  
Best Local Similarity 100.0%; Pred. No. 66;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGCTGCTGCACCTGT 16  
Db 67 GGCTGCTGCACCTGT 52

RESULT 22  
LOCUS AW732816/c 430 bp mRNA linear EST 21-APR-2000  
DEFINITION b14h10.y1 NIH\_MGC\_21 Homo sapiens cDNA clone IMAGE:2962915,  
similar to TR:O14801 O14801 EURARYOTIC TRANSLATION INITIATION  
FACTOR 3 SUBUNIT. [1] ; mRNA sequence.  
ACCESSION AW732816  
VERSION AW732816.1 GI:7633154  
KEYWORDS EST.  
SOURCE Homo sapiens (human)  
ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo;  
1 (bases 1 to 430)  
NIH-MGC http://mgi.nci.nih.gov/.  
National Institutes of Health, Mammalian Gene Collection (MGC)  
Unpublished  
Contact: Robert Strausberg, Ph.D.  
Email: cgabbs-r@mail.nih.gov  
Tissue Procurement: ATCC  
CDNA Library Preparation: Ling Hong/Rubin Laboratory  
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNL)  
DNA Sequencing by: Washington University Genome Sequencing Center  
Clone distribution: MGC clone distribution information can be  
found through the I.M.A.G.E. Consortium/LNL at:  
image.lnl.gov/image/html/resources.shtml  
Seq primer: -40RP from Gibco  
High quality sequence stop: 412.  
Location/Qualifiers

## FEATURES

SOURCE

1.430  
/organism="Homo sapiens"  
/mol\_type="mRNA"  
/db\_xref="taxon:9606"  
/clone="IMAGE:2962915"  
/issue\_type="choriocarcinoma"  
/lab\_host="DH10B (phage-resistant)"  
/clone\_lib="NIH MGC 21"  
/note="Organ: Placenta; Vector: pOT7; Site\_1: XhoI;  
Site\_2: EcoRI; cDNA made by oligo-dT priming.  
Directionally cloned into EcoRI/XhoI sites using the  
following 5' adaptor: GGCACGAG(G). Size-selected >500bp

for average insert size 1.8kb. Library constructed by  
Ling Hong in the laboratory of Gerald M. Rubin (University  
of California, Berkeley) using ZAP-cDNA synthesis kit  
(Stratagene) and Superscript II RT (Life Technologies)."  
BASE COUNT 119 a 112 c 125 g 74 t  
ORIGIN

Query Match 88.9%; Score 16; DB 9; Length 430;  
Best Local Similarity 100.0%; Pred. No. 66;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGCTGCTGCACCTGT 16  
Db 140 GGCTGCTGCACCTGT 125

RESULT 23  
LOCUS BF850584/c 431 bp mRNA linear EST 16-JAN-2001  
DEFINITION PML-EN0065-191100-007-h05 EN0065 Homo sapiens cDNA, mRNA sequence.  
ACCESSION BF850584  
VERSION BF850584.1 GI:12237746  
KEYWORDS EST.  
SOURCE Homo sapiens (human)  
ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo;  
1 (bases 1 to 431)  
Dias Neto, E., Garcia Correa, R., Verjovski-Almeida, S., Brines, M.R.,  
Nagai, M.A., da Silva, W. Jr., Zago, M.A., Bordin, S., Costa, F.F.,  
Goldman, G.H., Carvalho, A.F., Matsukuma, A., Bala, G.S., Simpson, D.H.,  
Brunstein, A., de Oliveira, P.S., Bucher, P., Jongeneel, C.V., O'Hare  
M.J., Soares, F., Brentani, R.R., Reis, L.F., de Souza, S.J. and  
Simpson, A.J.  
Shotgun sequencing of the human transcriptome with ORF expressed  
sequence tags  
Proc. Natl. Acad. Sci. U.S.A. 97 (7), 3491-3496 (2000)  
JOURNAL MEDLINE  
PUBMED 10737800  
CONTACT: Simpson A.J.G.  
Laboratory of Cancer Genetics  
Ludwig Institute for Cancer Research  
Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP,  
Brazil  
Tel: +55-11-2704932  
Fax: +55-11-2707001  
Email: asimpson@ludwig.org.br  
This sequence was derived from the FAPESP/LICR Human Cancer Genome  
Project. This entry can be seen in the following URL  
(http://www.ludwig.org.br/scripts/gethtml2.pl?tl=PML&t2=PML-EN0065-  
191100-007-h05&t3=2000-11-19&t4=1)  
Seq primer: puc 18 forward  
High quality sequence start: 39  
High quality sequence stop: 431.  
Location/Qualifiers

## FEATURES

SOURCE

1.431  
/organism="Homo sapiens"  
/mol\_type="mRNA"  
/db\_xref="taxon:9606"  
/dev\_stage="Adult"  
/clone\_lib="EN0065"  
/note="Organ: lung normal; Vector: puc18; Site\_1: SmaI;  
Site\_2: SmaI; A mini-library was made by cloning products  
derived from ORESTES PCR (O.S. Letters Patent application  
No. 196,716 - Ludwig Institute for Cancer Research)  
profiles into the pUC 18 vector. Reverse transcription of  
tissue mRNA and cDNA amplification were performed under  
low stringency conditions."  
BASE COUNT 65 a 157 c 107 g 102 t  
ORIGIN

Query Match 88.9%; Score 16; DB 10; Length 431;  
Best Local Similarity 100.0%; Pred. No. 66;

Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGCTGTCACCTGT 16  
 |||||  
 Db 361 GGCTGGTGCCTGT 346

RESULT 24  
 CB270361/c 431 bp mRNA linear EST 20-FEB-2003  
 LOCUS 1009268 Human Fat Cell 5'-Stretch Plus cDNA Library Homo sapiens  
 DEFINITION  
 CB270361  
 CDNA 5', mRNA sequence.  
 CB270361  
 CB270361.1 GI:28444946  
 EST.  
 SOURCE Homo sapiens (human)  
 ORGANISM Homo sapiens  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.  
 1 (bases 1 to 431)  
 Yang, R.-Z., Shuldiner, A. and Gong, D.-W.  
 EST analysis of human adipose gene expression  
 JOURNAL Unpublished  
 COMMENT Contact: Gong Da-Wei  
 Division of Endocrinology, Diabetes and Nutrition  
 University of Maryland  
 660 Redwood St. HH497, Baltimore, MD 21201, USA  
 Tel: 410 706 1672  
 Fax: 410 706 1622  
 Email: dgong@medicine.umaryland.edu  
 PCR Primers  
 FORWARD: CTCGGAGCGCGCATTTGTTGT  
 BACKWARD: AATGACTACTATGAGCGGATTTGG  
 Seq primer: GTTGTACCGCGGATTC.  
 Location/Qualifiers  
 1. 431  
 /organism="Homo sapiens"  
 /mol\_type="mRNA"  
 /db\_xref="taxon:9606"  
 /sex="Male and Female"  
 /tissue\_type="Adipose"  
 /clone\_lib="Human Fat Cell 5'-Stretch Plus cDNA library"  
 /note="Vector: lambdaTriplex"  
 112 c 125 g 74 t

BASE COUNT 120 a 112 c 125 g 74 t

ORIGIN

Query Match 88.9%; Score 16; DB 14; Length 431;  
 Best Local Similarity 100.0%; Pred. No. 66;  
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGCTGTCACCTGT 16  
 |||||  
 Db 135 GGCTGGTGCCTGT 120

RESULT 25  
 CB145604/c 433 bp mRNA linear EST 29-JAN-2003  
 LOCUS K-EST0200441 L11SNJ354s1 Homo sapiens CDNA clone L11SNJ354s1-13-C11  
 DEFINITION  
 CB145604  
 5', mRNA sequence.  
 CB145604  
 CB145604.1 GI:28125676  
 EST.  
 SOURCE Homo sapiens (human)  
 ORGANISM Homo sapiens  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.  
 1 (bases 1 to 433)  
 Kim, N.S., Hahn, Y., Oh, J.H., Lee, J.Y., Ahn, H.Y., Chu, M.Y., Kim, M.R.,  
 Oh, K.-J., Cheong, J.B., Sohn, H.Y., Kim, J.M., Park, H.S., Kim, S. and  
 Kim, Y.S.  
 21C Frontier Korean EST Project 2001  
 JOURNAL Unpublished

COMMENT Contact: Kim YS  
 Genome Research Center  
 Korea Research Institute of Bioscience & Biotechnology  
 52 Boeun-dong Yuseong-gu, Daejeon 305-333, South Korea  
 Tel: +82-42-860-4470  
 Fax: +82-42-860-4409  
 Email: yongsung@mail.kribb.re.kr  
 Plate: 13 row: C column: 11  
 High quality sequence stop: 433.  
 Location/Qualifiers  
 1. 433  
 /organism="Homo sapiens"  
 /mol\_type="mRNA"  
 /db\_xref="taxon:9606"  
 /clone="L11SNJ354s1-13-C11"  
 /sex="M"  
 /tissue\_type="Liver"  
 /cell\_type="Polygonal"  
 /cell\_line="SNU-354"  
 /lab\_host="Top10F"  
 /clone\_lib="L11SNJ354s1"  
 /note="Organ: Liver; Vector: PCNS-D2; Site 1: EcoRI;  
 Site 2: NotI. The poly (A) + RNA was dephosphorylated with  
 bacterial alkaline phosphatase (BAP) and then deacapped  
 with tobacco acid pyrophosphatase (TAP). The deacapped  
 intact mRNA was ligated with DNA-RNA linker including  
 EcoRI site by treatment of T4 RNA ligase and the first  
 strand cDNA was synthesized from oligo dt-selected mRNA by  
 priming with dt-tailed vector. The dt-tailed vector was  
 adjusted to have about 60nt. The cDNA vector was  
 circularized with E. coli DNA ligase after digestion of  
 EcoRI which site is also included in vector. An RNA strand  
 converted to a DNA strand by Okayama-Berg method. The  
 obtained cDNA vectors were used for transformation of  
 competent cells E. coli Top10F by electroporation method.  
 The cDNA libraries constructed by this method are  
 full-length enriched cDNA library. After analyzing and  
 sequencing about 2,000 - 3,000 colonies in original cDNA  
 library, the abundant cDNAs were selected and amplified by  
 PCR reaction using vector region primer including T7  
 promoter as 5' primer and Nidrl14 as 3' primer. The PCR  
 products were used as template for synthesis of  
 biotinylated single stranded RNA by in vitro transcription  
 reaction. The synthesized RNA probes were hybridized with  
 antisense single stranded cDNAs prepared from original  
 library and incubated with avidin-gel. After removing  
 DNA-RNA hybrids by centrifuge, the substracted cDNA  
 libraries were constructed by transformation of the  
 remaining DNA into competent cells E. coli Top10F with  
 electroporation method."  
 110 a 115 c 131 g 77 t

BASE COUNT 110 a 115 c 131 g 77 t

ORIGIN

Query Match 88.9%; Score 16; DB 14; Length 433;  
 Best Local Similarity 100.0%; Pred. No. 66;  
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGCTGTCACCTGT 16  
 |||||  
 Db 187 GGCTGGTGCCTGT 172

RESULT 26  
 CB129626/c 434 bp mRNA linear EST 29-JAN-2003  
 LOCUS K-EST019303 L11SNJ354 Homo sapiens CDNA clone L11SNJ354-16-B02 5',  
 DEFINITION  
 CB129626  
 mRNA sequence.  
 CB129626  
 CB129626.1 GI:28093354.  
 EST.  
 SOURCE Homo sapiens (human)  
 ORGANISM Homo sapiens  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

REFERENCE  
AUTHORS

Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.  
1 (bases 1 to 434)  
Kim,N.S., Hahn,Y., Oh,J.H., Lee,J.Y., Ahn,H.Y., Chu,M.Y., Kim,M.R.,  
Oh,K.J., Cheong,J.E., Sohn,H.Y., Kim,J.M., Park,H.S., Kim,S. and  
Kim,Y.S.

TITLE  
JOURNAL  
COMMENT

21C Frontier Korean EST Project 2001  
Unpublished  
Contact: Kim YS  
Genome Research Center  
Korea Research Institute of Bioscience & Biotechnology  
52 Boeun-dong Yuseong-gu, Daejeon 305-333, South Korea  
Tel: +82-42-860-4470  
Fax: +82-42-860-4409  
Email: yongsung@mail.kribb.re.kr  
Plate: 16 row: B column: 02  
High quality sequence stop: 434.

FEATURES  
source

1.434  
/organism="Homo sapiens"  
/mol\_type="mRNA"  
/db\_xref="taxon:9606"  
/clone="U11SNJ354-16-B02"  
/sex="M"  
/tissue\_type="liver"  
/cell\_type="Polygonal"  
/cell\_line="SNU-354"  
/lab\_host="Top10F"  
/clone\_11b="U11SNJ354"  
/note="Organ: Liver; Vector: pCNS-D2; Site: 1: EcoRI;  
Site: 2: NotI; The poly (A) + RNA was dephosphorylated with  
bacterial alkaline phosphatase (BAP) and then decapped  
with tobacco acid pyrophosphatase (TAP). The decapped  
intact mRNA was ligated with DNA-RNA linker including  
EcoRI site by treatment of T4 RNA ligase and the first  
strand cDNA was synthesized from oligo dt-selected mRNA by  
priming with dt-tailed vector. The dt-tailed vector was  
adjusted to have about 60nt. The cDNA vector was  
circularized with E. coli DNA ligase after digestion of  
EcoRI which site is also included in vector. An RNA strand  
converted to a DNA strand by Okayama-Berg method. The  
obtained cDNA vectors were used for transformation of  
competent cells E. coli Top10F by electroporation method.  
The cDNA libraries constructed by this method are  
full-length enriched cDNA library."

BASE COUNT 122 a 112 c 125 g 75 t  
ORIGIN

Query Match 88.9%; Score 16; DB 14; Length 434;  
Best Local Similarity 100.0%; Pred. No. 66;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGCTGCTGCACCTGT 16  
|||||  
Db 142 GGCTGCTGCACCTGT 127

RESULT 27 436 bp mRNA linear EST 29-JAN-2003  
CBI42213/C  
LOCUS  
DEFINITION  
K-EST0195953 U11SNJ354s1 Homo sapiens cDNA clone U11SNJ354s1-19-E11  
5', mRNA sequence.  
ACCESSION  
CBI42213  
VERSION  
CBI42213.1 GI:28118094  
KEYWORDS  
EST.  
SOURCE  
Homo sapiens (human)  
ORGANISM  
Homo sapiens

REFERENCE  
AUTHORS  
Kim,N.S., Hahn,Y., Oh,J.H., Lee,J.Y., Ahn,H.Y., Chu,M.Y., Kim,M.R.,  
Oh,K.J., Cheong,J.E., Sohn,H.Y., Kim,J.M., Park,H.S., Kim,S. and  
Kim,Y.S.  
TITLE  
21C Frontier Korean EST Project 2001

JOURNAL  
COMMENT

Unpublished  
Contact: Kim YS  
Genome Research Center  
Korea Research Institute of Bioscience & Biotechnology  
52 Boeun-dong Yuseong-gu, Daejeon 305-333, South Korea  
Tel: +82-42-860-4470  
Fax: +82-42-860-4409  
Email: yongsung@mail.kribb.re.kr  
Plate: 19 row: B column: 11  
High quality sequence stop: 436.  
Location/Qualifiers

FEATURES  
source

1.436  
/organism="Homo sapiens"  
/mol\_type="mRNA"  
/db\_xref="taxon:9606"  
/clone="U11SNJ354s1-19-E11"  
/sex="M"  
/tissue\_type="liver"  
/cell\_type="Polygonal"  
/cell\_line="SNU-354"  
/lab\_host="Top10F"  
/clone\_11b="U11SNJ354s1"  
/note="Organ: Liver; Vector: pCNS-D2; Site: 1: EcoRI;  
Site: 2: NotI; The poly (A) + RNA was dephosphorylated with  
bacterial alkaline phosphatase (BAP) and then decapped  
with tobacco acid pyrophosphatase (TAP). The decapped  
intact mRNA was ligated with DNA-RNA linker including  
EcoRI site by treatment of T4 RNA ligase and the first  
strand cDNA was synthesized from oligo dt-selected mRNA by  
priming with dt-tailed vector. The dt-tailed vector was  
adjusted to have about 60nt. The cDNA vector was  
circularized with E. coli DNA ligase after digestion of  
EcoRI which site is also included in vector. An RNA strand  
converted to a DNA strand by Okayama-Berg method. The  
obtained cDNA vectors were used for transformation of  
competent cells E. coli Top10F by electroporation method.  
The cDNA libraries constructed by this method are  
full-length enriched cDNA library. After analyzing and  
sequencing about 2,000 - 3,000 colonies in original cDNA  
library, the abundant cDNAs were selected and amplified by  
PCR reaction using vector region primer including T7  
promotor as 5' primer and NidT14 as 3' primer. The PCR  
products were used as template for synthesis of  
biotinylated single stranded RNA by in vitro transcription  
reaction. The synthesized RNA probes were hybridized with  
antisense single stranded cDNAs prepared from original  
library and incubated with avidin-gel. After removing  
DNA-RNA hybrids by centrifuge, the substracted cDNA  
libraries were constructed by transformation of the  
remaining DNA into competent cells E. coli Top10F with  
electroporation method."

BASE COUNT 112 a 113 c 132 g 79 t  
ORIGIN

Query Match 88.9%; Score 16; DB 14; Length 436;  
Best Local Similarity 100.0%; Pred. No. 66;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGCTGCTGCACCTGT 16  
|||||  
Db 172 GGCTGCTGCACCTGT 157

RESULT 28 438 bp mRNA linear EST 16-JAN-2001  
BF858511/C  
LOCUS  
DEFINITION  
R01-FT0190-221100-021-c02 F0190 Homo sapiens cDNA, mRNA sequence.  
ACCESSION  
BF858511  
VERSION  
BF858511.1 GI:12246255  
KEYWORDS  
EST.  
SOURCE  
Homo sapiens (human)  
ORGANISM  
Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

REFERENCE 1 (bases 1 to 438)  
AUTHORS Dias Neto,E., Garcia Correa,R., Verjovski-Almeida,S., Briones,M.R., Nagai,M.A., da Silva,M. Jr., Zago,M.A., Bordin,S., Costa,F.F., Goldman,G.H., Carvalho,A.F., Matsukuma,A., Baia,G.S., Simpson,D.H., Brunstein,A., deOliveira,P.S., Bucher,P., Jongeneel,C.V., O'Hare,M.J., Soares,F., Brentani,R.R., Reis,L.F., de Souza,S.J. and Simpson,A.J.  
TITLE Shotgun sequencing of the human transcriptome with ORF expressed sequence tags  
JOURNAL Proc. Natl. Acad. Sci. U.S.A. 97 (7), 3491-3496 (2000)  
MEDLINE 2020263  
PUBMED 10737800  
COMMENT Contact: Simpson A.J.G.  
Laboratory of Cancer Genetics  
Ludwig Institute for Cancer Research  
Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP, Brazil  
Tel: +55-11-2704922  
Fax: +55-11-2707001  
Email: asimpson@ludwig.org.br  
This sequence was derived from the FAPESP/LICR Human Cancer Genome Project. This entry can be seen in the following URL  
(http://www.ludwig.org.br/scripts/gethtml2.pl?l=RCL&t2=RCL-FT0190-221100-021-c02&t3=2000-11-22&t4=1)  
Seq primer: puc 18 forward  
High quality sequence start: 27  
Location/Qualifiers  
FEATURES  
source  
1. 438  
/organism="Homo sapiens"  
/mol\_type="mRNA"  
/db\_xref="taxon:9606"  
/dev\_stage="Adult"  
/clone\_1lb="FT0190"  
/note="Organ: prostate tumor; Vector: puc18; Site 1: SmaI; Site 2: SmaI; A mini-library was made by cloning products derived from ORESTES PCR (U.S. Letters Patent application No. 196,716 - Ludwig Institute for Cancer Research) profiles into the pUC 18 vector. Reverse transcription of tissue mRNA and cDNA amplification were performed under low stringency conditions."  
BASE COUNT 115 a 110 c 133 g 80 t  
ORIGIN  
Query Match 88.9%; Score 16; DB 10; Length 438;  
Best Local Similarity 100.0%; Pred. No. 66;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
Qy 1 GGCTGGTGCACCTGT 16  
119 GCGTGGTGCACCTGT 158  
Db 173 GCGTGGTGCACCTGT 158  
RESULT 29  
BG104409 447 bp mRNA linear EST 30-JAN-2001  
LOCUS 602311027F1 NIH\_MGC\_20 Homo sapiens cDNA clone IMAGE:4422918 5',  
DEFINITION mRNA sequence.  
ACCESSION BG104409  
VERSION BG104409.1 GI:12598251  
KEYWORDS EST.  
SOURCE Homo sapiens (human)  
ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.  
REFERENCE 1 (bases 1 to 447)  
NIH-MGC http://mgc.nci.nih.gov/.  
National Institutes of Health, Mammalian Gene Collection (MGC)  
JOURNAL Unpublished  
COMMENT Contact: Robert Strausberg, Ph.D.  
Email: cgabbs-r@mail.nih.gov  
Tissue Procurement: ATCC/DCTD/DTP

CDNA Library Preparation: Ling Hong/Rubin Laboratory  
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNL)  
DNA Sequencing by: Incyte Genomics, Inc.  
Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LNL at: image.lnl.gov  
Place: LNCMI218 row: 1 column: 07  
High quality sequence stop: 445.  
Location/Qualifiers  
FEATURES  
source  
1. 447  
/organism="Homo sapiens"  
/mol\_type="mRNA"  
/db\_xref="taxon:9606"  
/clone\_1lb="IMAGE:4422918"  
/tissue\_type="melanotic melanoma"  
/lab\_host="DH10B (phage-resistant)"  
/clone\_1lb="NIH MGC 20"  
/note="Organ: skin; Vector: pORF7; Site 1: XhoI; Site 2: EcoRI; CDNA made by oligo-dT priming. Directionally cloned into EcoRI/XhoI sites using the following 5' adaptor: GGCGAG(G). Size-selected >500bp for average insert size 1.8kb. Library constructed by Ling Hong in the laboratory of Gerald M. Rubin (University of California, Berkeley) using ZAP-cDNA synthesis kit (Stratagene) and Superscript II RT (Life Technologies)."  
BASE COUNT 129 a 116 c 134 g 68 t  
ORIGIN  
Query Match 88.9%; Score 16; DB 10; Length 447;  
Best Local Similarity 100.0%; Pred. No. 67;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
Qy 1 GGCTGGTGCACCTGT 16  
119 GCGTGGTGCACCTGT 104  
Db 119 GCGTGGTGCACCTGT 104  
RESULT 30  
BF130453 449 bp mRNA linear EST 24-OCT-2000  
LOCUS 601818761F1 NIH\_MGC\_58 Homo sapiens cDNA clone IMAGE:404398 5',  
DEFINITION mRNA sequence.  
ACCESSION BF130453  
VERSION BF130453.1 GI:10969493  
KEYWORDS EST.  
SOURCE Homo sapiens (human)  
ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.  
REFERENCE 1 (bases 1 to 449)  
NIH-MGC http://mgc.nci.nih.gov/.  
National Institutes of Health, Mammalian Gene Collection (MGC)  
JOURNAL Unpublished  
COMMENT Contact: Robert Strausberg, Ph.D.  
Email: cgabbs-r@mail.nih.gov  
Tissue Procurement: ATCC  
CDNA Library Preparation: CLONETECH Laboratories, Inc.  
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNL)  
DNA Sequencing by: Incyte Genomics, Inc.  
Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LNL at:  
http://image.lnl.gov  
Plate: LNCM868 row: 3 column: 15  
High quality sequence stop: 449.  
Location/Qualifiers  
FEATURES  
source  
1. 449  
/organism="Homo sapiens"  
/mol\_type="mRNA"  
/db\_xref="taxon:9606"  
/clone\_1lb="IMAGE:404398"  
/tissue\_type="hypernephroma"  
/lab\_host="DH10B (T1 phage-resistant)"  
/clone\_1lb="NIH\_MGC\_58"  
/note="Organ: Kidney; Vector: pDNR-LIB (Clontech); Site 1:

Sf11 (ggcgcgtcgcgc); Site 2: Sf11 (ggccattatggcc);  
Double-stranded cDNA was prepared from cell line RNA. 5' and 3' adaptors were used in cloning as follows: 5' adaptor sequence: 5'-CACGGCCATTATGACC-3' and 3' adaptor sequence: 5'-ATTCTAGAGCGGAGCGCCGACATG-dT(30)BN-3' (where B = A, C, G or T). Average insert size 1.35 kb (range 0.9-4.0 kb). 15/15 colonies contained inserts by PCR. This library was enriched for full-length clones and was constructed by Clontech Laboratories (Palo Alto, CA)."

BASE COUNT 110 a 124 c 130 g 85 t

ORIGIN

Query Match 88.9%; Score 16; DB 10; Length 449;  
Best Local Similarity 100.0%; Pred. No. 67;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 GGCTGGTGCACCTGT 16  
122 GGCTGGTGCACCTGT 107

Db

RESULT 31  
BE909128 452 bp mRNA linear EST 20-OCT-2000  
LOCUS BE909128/c  
DEFINITION BE909128F1 NIH\_MGC\_70 Homo sapiens cDNA clone IMAGE:3903510 5', mRNA sequence.

ACCESSION BE909128  
VERSION BE909128.1 GI:10404401  
KEYWORDS EST.  
SOURCE Homo sapiens (human)  
ORGANISM Homo sapiens

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo. 1 (bases 1 to 452)  
AUTHORS NIH-MGC <http://mgs.nci.nih.gov/>.  
TITLE National Institutes of Health, Mammalian Gene Collection (MGC)  
JOURNAL Unpublished  
COMMENT Contact: Robert Strausberg, Ph.D.  
Email: [cgabbs-r@mail.nih.gov](mailto:cgabbs-r@mail.nih.gov)  
Tissue Procurement: ATCC  
cDNA Library Preparation: Life Technologies, Inc.  
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)  
DNA Sequencing by: Incyte Genomics, Inc.  
Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: <http://image.llnl.gov>  
Plate: L1AM9707 row: 1 column: 07  
High quality sequence stop: 452.  
Location/Qualifiers

FEATURES  
source 1..452  
/organism="Homo sapiens"  
/mol\_type="mRNA"  
/db\_xref="taxon:9606"  
/clone="IMAGE:3903510"  
/tissue\_type="epithelioid carcinoma"  
/lab\_host="DH10B (phage-resistant)"  
/clone\_11b="NIH\_MGC\_70"  
/note="Organ: pancreas; Vector: PCMV-SPORT6; Site 1: NCI; Site 2: Sall; Cloned unidirectionally. Primer: Oligo dT. Average insert size 1.1 kb. Library constructed by Life Technologies."

BASE COUNT 127 a 117 c 133 g 75 t

ORIGIN

Query Match 88.9%; Score 16; DB 10; Length 452;  
Best Local Similarity 100.0%; Pred. No. 67;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 GGCTGGTGCACCTGT 16  
137 GGCTGGTGCACCTGT 122

Db

RESULT 32  
BX475327/c 452 bp mRNA linear EST 12-JUN-2003  
LOCUS BX475327/c  
DEFINITION DKFZp686109176.r1 686 (synonym: h1cc3) Homo sapiens cDNA clone  
ACCESSION DKFZp686109176.5, mRNA sequence.  
VERSION BX475327  
KEYWORDS BX475327.1 GI:31672610  
EST.

SOURCE Homo sapiens (human)

ORGANISM Homo sapiens

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo. 1 (bases 1 to 452)

AUTHORS Bahr, A., Lauber, J., Mewes, H.W., Weil, B., Amd, C., Osanger, A., Fobo, G., Han, M., and Wiemann, S.  
TITLE (Bahr, A., Lauber, J., Mewes, H.W., Weil, B., et al.)  
JOURNAL Unpublished  
COMMENT Contact: Bahr A  
MIPS

Ingolstaedter Landstr.1, D-85764 Neuberg, Germany  
This is the 5' sequence of the clone insert  
Clone from S. Wiemann, Molecular Genome Analysis, German Cancer Research Center (DKFZ); Email [s.wiemann@dkfz-heidelberg.de](mailto:s.wiemann@dkfz-heidelberg.de); sequenced by Qiagen (Hilden/Germany) within the cDNA sequencing consortium of the German Genome Project.  
No sl sequence available.  
This clone (DKFZp686109176) is available at the RZPD in Berlin. Please contact the RZPD: Ressourcenzentrum, Heubnerweg 6, 14059 Berlin-Charlottenburg, GERMANY; Email: [clone@rzpd.de](mailto:clone@rzpd.de).

FEATURES  
source 1..452  
Location/Qualifiers

/organism="Homo sapiens"  
/mol\_type="mRNA"  
/db\_xref="taxon:9606"  
/clone="DKFZp686109176"  
/tissue\_type="human skeletal muscle"  
/dev\_host="radult"  
/lab\_host="DH10B"  
/clone\_11b="686 (synonym: h1cc3)"  
/note="Vector: pTRIPLEX2; Site\_1: Sf11A; Site\_2: Sf11B; cDNA-collection"

BASE COUNT 114 a 120 c 138 g 80 t

ORIGIN

Query Match 88.9%; Score 16; DB 13; Length 452;  
Best Local Similarity 100.0%; Pred. No. 67;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 GGCTGGTGCACCTGT 16  
184 GGCTGGTGCACCTGT 169

Db

RESULT 33  
AA496839/c 453 bp mRNA linear EST 12-AUG-1997  
LOCUS AA496839  
DEFINITION a633b02.r1 Gessler Wilms tumor Homo sapiens cDNA clone IMAGE:897579  
5' similar to contains Alu repetitive element; contains element TAR1 repetitive element; mRNA sequence.

ACCESSION AA496839  
VERSION AA496839.1 GI:2230160  
KEYWORDS EST.

SOURCE Homo sapiens (human)

ORGANISM Homo sapiens

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo. 1 (bases 1 to 453)

AUTHORS Hillier, L., Allen, M., Bowles, L., Dubuque, T., Geisel, G., Jost, S., Kucaba, T., Lacy, M., Le, N., Lennon, G., Marra, M., Martin, J., Moore, B., Scheinberg, K., Stepcie, M., Tan, F., Theising, B., White, Y., Wyllie, T., Waterston, R., and Wilson, R.  
TITLE WashU-Merck EST Project 1997

JOURNAL  
COMMENT

Unpublished  
Contact: Wilson RK  
Washington University School of Medicine  
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108  
Tel: 314 286 1800  
Fax: 314 286 1810  
Email: est@wustl.wustl.edu  
This clone is available royalty-free through LNLN; contact the  
IMAGE Consortium (info@image.llnl.gov) for further information.  
High quality sequence stop: 427.

FEATURES  
source

1.453  
/organism="Homo sapiens"  
/mol\_type="mRNA"  
/db\_xref="taxon:9606"  
/clone="IMAGE:897579"  
/sex="pooled (6)"  
/lab\_host="DH10B"  
/clone\_lib="Gessler Wilms tumor"  
/note="Vector: pSPORT1, Site 1: SalI; Site 2: NotI; RNA  
was prepared from a pool of 6 anonymous Wilms' tumor RNAs.  
RNA was prepared by acid-phenol, followed by one round of  
oligo dT selection. cDNA library preparation was with  
the BRL/Life Tech. Superscript Plasmid system. An  
oligo-dT NotI primer for first strand synthesis generated  
ggcgccgcc(t)n at the 3' end of the clones. A 5' SalI  
adaptor was used with sequence 5'-gtcagccagcgcg-3'.  
Resulting cDNAs were size selected (average size 2 kb),  
NotI digested, and ligated into NotI/SalI-cut pSPORT1.  
Library was constructed by Dr. Manfred Gessler."

BASE COUNT  
ORIGIN

115 a 124 c 135 g 79 t

Query Match 88.9%; Score 16; DB 9; Length 453;  
Best Local Similarity 100.0%; Pred. No. 67;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GGCTGCTGCACCTGT 16  
|||||  
Db 117 GGCTGCTGCACCTGT 102

RESULT 34  
BM171994 454 bp mRNA linear EST 04-DEC-2001  
LOCUS imageg3.3 2001/mm196bdf41.xl NIH\_MGC\_61 Homo sapiens cDNA clone  
DEFINITION IMAGE:468687 5', mRNA sequence.

ACCESSION BM171994  
VERSION BM171994.1 GI:17311557  
KEYWORDS EST.  
SOURCE Homo sapiens (human)  
ORGANISM Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE 1 (bases 1 to 454)  
AUTHORS Kale, P.T., Harsch, T.J., Folta, P.A., Nelson, D.O., Sanders, C.G. and  
Prange, C.K.

TITLE The I.M.A.G.E. Consortium quality control effort: clone  
resequencing for verification  
JOURNAL Unpublished  
COMMENT Other\_ESTS: BG529561  
Contact: Prange CK  
The I.M.A.G.E. Consortium  
Lawrence Livermore National Laboratory  
Livermore, CA, USA  
Email: help@image.llnl.gov

This read has been verified (found to hit its original self in the  
correct orientation), as part of the I.M.A.G.E. Consortium quality  
control effort. High quality sequence is defined as having 100 or  
more base pairs with a paired quality value of 20 or greater, where  
a sliding window of 4 base pairs with a paired quality value of 15  
or greater marks the beginning and end of the sequence. For  
information on obtaining this clone, please contact

info@image.llnl.gov.  
Plate: L1CM1498 row: k column: 18  
Seq primer: -21m13  
High quality sequence stop: 454.  
Location/Qualifiers

FEATURES  
source

1.454  
/organism="Homo sapiens"  
/mol\_type="mRNA"  
/db\_xref="taxon:9606"  
/clone="IMAGE:468687"  
/tissue\_type="embryonal carcinoma"  
/lab\_host="DH10B (T1 phage-resistant)"  
/clone\_lib="NIH\_MGC\_61"  
/note="Organ: testis; Vector: pDNR-LIB (Clontech); Site 1:  
SfiI (ggcgccgcgcgc); Site 2: SfiI (ggcgccatcgcc);  
Double-stranded cDNA was prepared from cell line RNA. 5'  
and 3' adaptors were used in cloning as follows: 5'  
adaptor sequence: 5'-ATCTAGAGCGCGAGCGCGCGACATG-dT(30)BN-3'  
(where B = A, C, G, or T). Average  
insert size 1.75 kb (range 0.9-4.0 kb). 15/15 colonies  
contained inserts by PCR. This library was enriched for  
full-length clones and was constructed by Clontech  
Laboratories (Palo Alto, CA). Note: this is a NIH\_MGC  
Library."

BASE COUNT  
ORIGIN

74 a 146 c 117 g 116 t 1 others

Query Match 88.9%; Score 16; DB 12; Length 454;  
Best Local Similarity 100.0%; Pred. No. 67;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GGCTGCTGCACCTGT 16  
|||||  
Db 208 GGCTGCTGCACCTGT 223

RESULT 35  
BF035036 455 bp mRNA linear EST 20-OCT-2000  
LOCUS 601456130F1 NIH\_MGC\_66 Homo sapiens cDNA clone IMAGE:3859734 5',  
DEFINITION mRNA sequence.

ACCESSION BF035036  
VERSION BF035036.1 GI:10742748  
KEYWORDS EST.  
SOURCE Homo sapiens (human)  
ORGANISM Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE 1 (bases 1 to 455)  
AUTHORS NIH-MGC http://mgs.nci.nih.gov/.  
TITLES National Institutes of Health, Mammalian Gene Collection (MGC)  
JOURNAL Unpublished  
COMMENT Contact: Robert Strausberg, Ph.D.  
Email: cgabbs-remail.nih.gov  
Tissue Procurement: DCTD/DTF  
cDNA library Preparation: Life Technologies, Inc.  
DNA Sequencing by: Incyte Genomics, Inc.  
Clone distribution: MGC clone distribution information can be  
found through the I.M.A.G.E. Consortium/LLNL at:  
http://image.llnl.gov  
Plate: L1AM9593 row: 1 column: 07  
High quality sequence stop: 455.

FEATURES  
source

1.455  
/organism="Homo sapiens"  
/mol\_type="mRNA"  
/db\_xref="taxon:9606"  
/clone="IMAGE:3859734"  
/tissue\_type="adenocarcinoma"  
/lab\_host="DH10B (phage-resistant)"  
/clone\_lib="NIH\_MGC\_66"

/note="Organ: ovary; Vector: pCMV-SPORT6; Site\_1: NotI;  
 Site\_2: SalI; Cloned unidirectionally. Primer: Oligo dt.  
 Average insert size 1.8 kb. Library constructed by Life  
 Technologies."  
 BASE COUNT 129 a 117 c 133 g 76 t  
 ORIGIN

Query Match 88.9%; Score 16; DB 10; Length 455;  
 Best Local Similarity 100.0%; Pred. No. 67;  
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 GGCTGGTGTACCTGT 16  
 |||||  
 Db 138 GGCTGGTGTACCTGT 123

Search completed: August 15, 2003, 10:57:39  
 Job time : 1136.8 secs .

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GenCore version 5.1.6  
Copyright (c) 1993 - 2003 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: August 14, 2003, 21:32:41 ; Search time 492.975 Seconds  
(without alignments)  
1493.734 Million cell updates/sec

Title: US-10-074-620-1.g  
Perfect score: 18  
Sequence: 1 ggcgtgctgcacctgtta 18

Scoring table: OLIGO\_NUC  
Gapop 60.0 , Gapext 60.0

Searched: 2888711 seqs, 2045481336 residues

Word size : 0

Total number of hits satisfying chosen parameters: 5777422

Minimum DB seq length: 0  
Maximum DB seq length: 200000000

Post-processing: Listing first 120 summaries

Database :

GenBank1: \*  
1: gb\_ba: \*  
2: gb\_hgt: \*  
3: gb\_in: \*  
4: gb\_om: \*  
5: gb\_ov: \*  
6: gb\_ph: \*  
7: gb\_pat: \*  
8: gb\_pl: \*  
9: gb\_pr: \*  
10: gb\_ro: \*  
11: gb\_sts: \*  
12: gb\_sy: \*  
13: gb\_un: \*  
14: gb\_vi: \*  
15: em\_ba: \*  
16: em\_fm: \*  
17: em\_hum: \*  
18: em\_in: \*  
19: em\_mu: \*  
20: em\_om: \*  
21: em\_or: \*  
22: em\_ov: \*  
23: em\_pat: \*  
24: em\_ph: \*  
25: em\_pl: \*  
26: em\_ro: \*  
27: em\_sts: \*  
28: em\_un: \*  
29: em\_vi: \*  
30: em\_hgt\_hum: \*  
31: em\_hgt\_inv: \*  
32: em\_hgt\_other: \*  
33: em\_hgt\_mus: \*  
34: em\_hgt\_pln: \*  
35: em\_hgt\_rtd: \*  
36: em\_hgt\_man: \*  
37: em\_hgt\_vrt: \*  
38: em\_sy: \*  
39: em\_hgt\_hum: \*  
40: em\_hgt\_mus: \*  
41: em\_hgt\_other: \*

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,  
and is derived by analysis of the total score distribution.

# SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	18	100.0	18	AX522236	AX522236 Sequence
2	18	100.0	175905	AC137657	AC137657 Sus scrofa
3	17	94.4	88557	AL136314	AL136314 Human DNA
4	17	94.4	107979	AP006146	AP006146 Lotus jap
5	17	94.4	226979	AC128140	AC128140 Rattus no
6	17	94.4	231014	AC096132	AC096132 Rattus no
7	17	94.4	238552	AC140198	AC140198 Mus muscu
8	17	94.4	244424	AC131614	AC131614 Rattus no
9	17	94.4	244636	AC123215	AC123215 Rattus no
10	17	94.4	248793	AC103080	AC103080 Rattus no
11	16	88.9	20	126102	126102 Sequence 28
12	16	88.9	50	AR032934	AR032934 Sequence
13	16	88.9	50	AR209598	AR209598 Sequence
14	16	88.9	50	129674	129674 Sequence 54
15	16	88.9	50	191348	191348 Sequence 54
16	16	88.9	435	BD025521	BD025521 Sequence
17	16	88.9	455	AX321046	AX321046 Sequence
18	16	88.9	963	BT006889	BT006889 Homo sapi
19	16	88.9	963	BT007572	BT007572 Synthetic
20	16	88.9	1041	104174	104174 Sequence 3
21	16	88.9	1103	AF020833	AF020833 Homo sapi
22	16	88.9	1115	HS096074	HS096074 Human trans
23	16	88.9	1128	BC000733	BC000733 Homo sapi
24	16	88.9	1138	BC008469	BC008469 Homo sapi
25	16	88.9	1142	BD063236	BD063236 Secreted
26	16	88.9	1174	AF094850	AF094850 Homo sapi
27	16	88.9	2199	AF348456	AF348456 Desmodem
28	16	88.9	2556	AF172333	AF172333 Human her
29	16	88.9	2661	AF172332	AF172332 Human her
30	16	88.9	2661	14 EBVBLFL1A	EBVBLFL1A
31	16	88.9	2663	14 EBVBLFL1	EBVBLFL1
32	16	88.9	2721	6 E00513	E00513 Genomic DNA
33	16	88.9	3210	14 HS4GP340A	HS4GP340A
34	16	88.9	3400	6 E01006	E01006 CDNA encodi
35	16	88.9	3400	6 E01006	E01006 CDNA encodi
36	16	88.9	3833	6 AR049357	AR049357 Sequence
37	16	88.9	4988	14 HS4GP340B	HS4GP340B
38	16	88.9	5019	6 H11178	H11178 Synthetic n
39	16	88.9	5019	6 E01007	E01007 DNA sequenc
40	16	88.9	5931	14 AR233080	AR233080 Sequence
41	16	88.9	5931	14 HS4ENYGP	HS4ENYGP
42	16	88.9	6450	9 AF092453	AF092453 Homo sapi
43	16	88.9	44904	5 TRUS04410	TRUS04410
44	16	88.9	108036	9 AC084352	AC084352 Homo sapi
45	16	88.9	128641	10 AL845521	AL845521 Mouse DNA
46	16	88.9	129048	9 AC020931	AC020931 Homo sapi
47	16	88.9	168012	2 AC025565	AC025565 Homo sapi
48	16	88.9	169734	9 AC106748	AC106748 Homo sapi
49	16	88.9	170803	2 AC016155	AC016155 Homo sapi
50	16	88.9	171823	14 HNV507799	HNV507799 Human her
51	16	88.9	172281	14 EBV	EBV
52	16	88.9	173950	9 AL390961	AL390961 Human DNA
53	16	88.9	174937	2 AC022245	AC022245 Homo sapi
54	16	88.9	184113	14 HS4B958RMJ	HS4B958RMJ
55	16	88.9	194148	2 AC125349	AC125349 Mus muscu
56	16	88.9	196360	2 AC125349	AC125349 Mus muscu
57	16	88.9	217717	2 BX530016	BX530016 Mus muscu
58	16	88.9	225024	2 AC116415	AC116415 Mus muscu
59	16	88.9	234288	2 AL772311	AL772311 Mus muscu
60	16	88.9	244595	2 AC095144	AC095144 Rattus no
61	16	88.9	244784	2 AC117356	AC117356 Rattus no
62	16	88.9	247249	2 AC023526	AC023526 Homo sapi
63	16	88.9	266738	2 AL138896	AL138896 Homo sapi
64	15	83.3	266	11 G37013	G37013 SHGC-56377
65	15	83.3	622	11 BV025210	BV025210 S212P6604

C	66	15	83.3	688	10	AF327854	AF327854	Spermophi
	67	15	83.3	781	11	BV059337	BV059337	S212P6928
	68	15	83.3	1419	6	E54455	E54455	Gene causat
	69	15	83.3	1419	10	AF124044	AF124044	Mus muscu
	70	15	83.3	1808	6	AK262598	AK262598	Sequence
	71	15	83.3	2293	6	E30803	E30803	Novel prote
	72	15	83.3	2352	6	E30804	E30804	Novel prote
	73	15	83.3	2951	6	AX017503	AX017503	Sequence
	74	15	83.3	2952	6	BD135193	BD135193	Human nuc
	75	15	83.3	3034	10	AK122327	AK122327	Mus muscu
	76	15	83.3	3065	6	AK262597	AK262597	Sequence
	77	15	83.3	3233	3	TBR308995	TBR308995	Typanoso
	78	15	83.3	3306	10	AB017609	AB017609	Mus muscu
	79	15	83.3	3337	6	E30802	E30802	Novel prote
	80	15	83.3	3660	10	AB017608	AB017608	Mus muscu
	81	15	83.3	3674	6	E30801	E30801	Novel prote
	82	15	83.3	3680	10	BC017126	BC017126	Mus muscu
	83	15	83.3	4190	10	M60103	M60103	Rattus norv
	84	15	83.3	4685	10	RNLMAR1	RNLMAR1	R. norvegicu
	85	15	83.3	6412	9	BC048768	BC048768	Homo sapi
	86	15	83.3	6545	10	RATLARA	RATLARA	Sequence
	87	15	83.3	6550	6	AX453123	AX453123	Sequence
	88	15	83.3	6553	6	AX453171	AX453171	Sequence
	89	15	83.3	6553	6	AF332878	AF332878	Cochliobo
	90	15	83.3	6672	6	AX335949	AX335949	Sequence
	91	15	83.3	6672	6	HSU18259	HSU18259	Human clone
	92	15	83.3	7702	6	AX658135	AX658135	Sequence
	93	15	83.3	7702	9	HSLARR	HSLARR	Human mRNA
	94	15	83.3	10040	10	AF467766	AF467766	Mus muscu
	95	15	83.3	14454	10	AB019041	AB019041	Mus muscu
	96	15	83.3	33608	10	AC003057	AC003057	Mouse Cos
	97	15	83.3	40438	10	AC087410	AC087410	Homo sapi
	98	15	83.3	42115	6	AX453180	AX453180	Sequence
	99	15	83.3	43101	3	AY198941	AY198941	Drosophi1
	100	15	83.3	43927	9	HSU69568	HSU69568	Human K28
	101	15	83.3	51730	9	AF324889S2	AF324889	Homo sapi
	102	15	83.3	53234	9	AC130889	AC130889	Homo sapi
	103	15	83.3	57224	2	AC136542	AC136542	Rattus no
	104	15	83.3	60761	2	AC102704	AC102704	Mus muscu
	105	15	83.3	67197	9	AF510423S1	AF510423	Homo sapi
	106	15	83.3	71983	9	AL445429	AL445429	Human DNA
	107	15	83.3	72000	9	AP001102	AP001102	Homo sapi
	108	15	83.3	73597	2	AL109598	AL109598	Homo sapi
	109	15	83.3	74089	9	AL589762	AL589762	Human DNA
	110	15	83.3	82419	9	AC004979	AC004979	Homo sapi
	111	15	83.3	83549	9	AC004752	AC004752	Homo sapi
	112	15	83.3	87157	9	AL136143	AL136143	Human DNA
	113	15	83.3	88376	9	AL514407	AL514407	Human DNA
	114	15	83.3	88554	10	AL669960	AL669960	Mouse DNA
	115	15	83.3	90497	9	HS732E4	HS732E4	Human DNA
	116	15	83.3	101882	9	AC021089	AC021089	Homo sapi
	117	15	83.3	101956	2	AP002494	AP002494	Homo sapi
	118	15	83.3	108765	2	AF216674	AF216674	Homo sapi
	119	15	83.3	109391	9	AL160031	AL160031	Human DNA
	120	15	83.3	110000	2	AC106541_3	AC106541	Continuation (4 of

## ALIGNMENTS

RESULT 1  
LOCUS AX522236 18 bp DNA  
DEFINITION Sequence 1 from Patent WO02064842.  
ACCESSION AX522236  
VERSION AX522236.1 GI:24411114  
KEYWORDS  
SOURCE Human herpesvirus 4 (Epstein-Barr virus)  
ORGANISM Human herpesvirus 4  
VIRUSES dsDNA viruses, no RNA stage; Herpesviridae;  
Gammaherpesvirinae; Lymphocryptovirus.

REFERENCE  
AUTHORS Wille, D. P. and Groen, P. A.

TITLE Quantitative epstein barr virus pcr rapid assay  
JOURNAL Patent: WO 02064842-A 1 22-AUG-2002;  
CHILDREN's Hospital Research Foundation (US)  
LOCATION/Qualifiers  
FEATURES  
Source  
1. 18  
/organism="Human herpesvirus 4"  
/mol\_type="genomic DNA"  
/db\_xref="taxon:10376"  
BASE COUNT  
2 a 4 c 6 g 6 t  
ORIGIN

Query Match 100.0%; Score 18; DB 6; Length 18;  
Best Local Similarity 100.0%; Pred. No. 0.81;  
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGCTGGTGCACCTGTTA 18  
DB 1 GGCTGGTGCACCTGTTA 18

RESULT 2  
LOCUS AC137657/c 175005 bp DNA linear HTG 21-MAR-2003  
DEFINITION Sus scrofa clone RP44-494H16, WORKING DRAFT SEQUENCE, 12 ordered  
pieces.  
ACCESSION AC137657  
VERSION AC137657.2 GI:29135576  
KEYWORDS HTG; HTGS\_PHASE2; HTGS\_DRAFT.  
SOURCE Sus scrofa (pig)  
ORGANISM Sus scrofa

REFERENCE  
AUTHORS Bukaryoga; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Cetartiodactyla; Suina; Suidae; Sus.  
1 (bases 1 to 175005)

Akher, N., Antonelli, A., Ayele, K., Beckett, S., Sternberg, S.M.,  
Benjamin, B., Blakesley, R.W., Bouffard, G.G., Brinkley, C., Brooks, S.,  
Carriaga, K., Coleman, B., Engle, J., Granite, S., Guan, X., Gupta, J.,  
Haghighi, P., Han, J., Hansen, N., Ho, S.-L., Idol, J.R., Karlins, E.,  
Laric, P., Lee-Lin, S.-Q., Legaspi, R., Maduro, Q.L., Maduro, V.B.,  
Margulies, E.H., Masello, C., Maskeri, B., Mcowell, J.,  
Paguirian, C., Pearson, R., Portnoy, M.E., Prasad, A.,  
Reddy-Bugue, N., Schandier, K., Scheller, M.G., Sison, C.,  
Stankiv, S., Thomas, J.W., Thomas, P.J., Touchman, J.W., Vogt, J.L.,  
Wetherby, K.D., Wiggins, L., Young, A., and Green, E.D.  
NISC Comparative Sequencing Initiative

TITLE Unpublished  
JOURNAL 2 (bases 1 to 175005)

REFERENCE  
AUTHORS Green, E.D.  
JOURNAL Direct Submission  
SUBMITTED (27-NOV-2002) NIH Intramural Sequencing Center, 8717  
Grovermont Circle, Gaithersburg, MD 20877, USA  
3 (bases 1 to 175005)

REFERENCE  
AUTHORS Green, E.D.  
JOURNAL Direct Submission  
SUBMITTED (21-MAR-2003) NIH Intramural Sequencing Center, 8717  
Grovermont Circle, Gaithersburg, MD 20877, USA  
On Mar 21, 2003 this sequence version replaced gi:25700398.

COMMENT ----- Genome Center

Center: NIH Intramural Sequencing Center  
Center code: NISC  
Web site: <http://www.nisc.nih.gov>  
Contact: [nisc.zoo@nhgri.nih.gov](mailto:nisc.zoo@nhgri.nih.gov)  
----- Project Information  
Center project name: ecg  
Center clone name: 494H16

The sequence data in this record represents an 'enhanced' version of a Phase 2 submission. Specifically, the indicated order and orientation of each sequence contig has been established using one or more of the following: read-pair data from individual subclones, overlaps with neighboring clones, alignment with available reference sequence (e.g., human), and/or confirmation by PCR testing. In addition, the sequence assembly is based on at least 8x average

coverage in Q20 bases and has been reviewed to rule out gross misassemblies, the low-quality ends of sequence contigs have been trimmed away, and each base is associated with a Phrap-derived quality score.

#### ----- Summary Statistics -----

Sequencing vector: plasmid; n/a; 100% of reads  
 Chemistry: Dye-terminator Big Dye; 100% of reads  
 Assembly program: Phrap; version 0.990319  
 Consensus quality: 171640 bases at least Q40  
 Consensus quality: 172768 bases at least Q30  
 Consensus quality: 173465 bases at least Q20  
 Insert size: 168000; agarose-IP  
 Quality coverage: 9.32x in Q20 bases; agarose-IP  
 Quality coverage: 9.01x in Q20 bases; sum-of-contigs

NOTE: This is a 'working draft' sequence. It currently consists of 12 contigs. Gaps between the contigs are represented as runs of N. The order of the pieces is believed to be correct as given, however the sizes of the gaps between them are based on estimates that have been provided by the submitter.

This sequence will be replaced by the finished sequence as soon as it is available and the accession number will be preserved.

```

1      8074: contig of 8074 bp in length
*      8075      8174: gap of unknown length
*      8175      33811: contig of 25637 bp in length
*      33812      33911: gap of unknown length
*      33912      36776: contig of 2765 bp in length
*      36777      36776: gap of unknown length
*      36777      50073: contig of 13297 bp in length
*      50074      50173: gap of unknown length
*      50174      57164: contig of 6991 bp in length
*      57165      57264: gap of unknown length
*      57265      59445: contig of 2181 bp in length
*      59446      59445: gap of unknown length
*      59446      72773: contig of 13228 bp in length
*      72774      72873: gap of unknown length
*      72874      124522: contig of 51649 bp in length
*      124523      124622: gap of unknown length
*      124623      143805: contig of 19183 bp in length
*      143806      143905: gap of unknown length
*      143906      149152: contig of 5247 bp in length
*      149153      149252: gap of unknown length
*      149253      167607: contig of 18355 bp in length
*      167608      167708: gap of unknown length
*      167708      175005: contig of 7298 bp in length.

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#### FEATURES

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  /organism="Sus scrofa"
  /mol_type="genomic DNA"
  /db_xref="taxon:9823"
  /clone="RP4-494H16"
  /clone_1fb="RP4"
  1..8074
    /note="assembly_fragment"
    clone_end:17
    vector_side:left
  8175..33811
    /note="assembly_fragment"
    33912..36676
      /note="assembly_fragment"
    36777..50073
      /note="assembly_fragment"
    50174..57164
      /note="assembly_fragment"
    57265..59445
      /note="assembly_fragment"
    59546..72773
      /note="assembly_fragment"
    72874..124522
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misc_feature 124623..143805
              /note="assembly_fragment"
misc_feature 143906..149152
              /note="assembly_fragment"
misc_feature 149253..167607
              /note="assembly_fragment"
misc_feature 167708..175005
              /note="assembly_fragment"

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BASE COUNT 36244 a 47550 c 50242 g 39869 t 1100 others
ORIGIN

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Query Match 100.0%; Score 18; DB 2; Length 175005;
Best Local Similarity 100.0%; Pred. No. 0.66;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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```

QY 1 GGCTGGTGCACCTGTTA 18
Db 18124 GGCTGGTGCACCTGTTA 18107

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```

RESULT 3
AL136314/c 88557 bp DNA linear PRI 01-NOV-2000
LOCUS Human DNA sequence from clone RP4-579A14 on chromosome 6, complete
DEFINITION

```

```

ACCESSION AL136314
VERSION AL136314.12 GI:11120979
KEYWORDS HTG.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens

```

```

REFERENCE Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
AUTHORS Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
TITLE 1 (bases 1 to 88557)
JOURNAL Peck,A.

```

#### COMMENT

Submitted (31-OCT-2000) Sanger Centre, Hinxton, Cambridgeshire, CB10 1SA, UK. E-mail enquiries: humquerry@sanger.ac.uk  
 requests: clonerequest@sanger.ac.uk  
 On Nov 8, 2000 this sequence version replaced gi:10931919.  
 During sequence assembly data is compared from overlapping clones. Where differences are found these are annotated as variations together with a note of the overlapping clone name. Note that the variation annotation may not be found in the sequence submission corresponding to the overlapping clone, as we submit sequences with only a small overlap as described above.  
 This sequence has been finished according to sequence map criteria as follows. An attempt is made to resolve all sequencing problems, such as compressions and repeats, but not necessarily within known annotated human repeat sequence elements (e.g. Alu). Where the sequence is ambiguous, there is an annotation using the 'unsure' feature key.  
 The following abbreviations are used to associate primary accession numbers given in the feature table with their source databases:  
 Em: EMBL; Sw: SWISSPROT; Tr: TREMBL; Wp: WORMPEP; Information on the WORMPEP database can be found at  
 http://www.sanger.ac.uk/Projects/C\_elegans/wormpep This sequence was generated from part of bacterial clone contigs of human chromosome 6, constructed by the Sanger Centre Chromosome 6 Mapping Group. Further information can be found at  
 http://www.sanger.ac.uk/HGP/Chr6  
 RP4-579A14 is from the library RPC1-4 constructed at the Roswell Park Cancer Institute by the group of Pieter de Jong. For further details see http://bacpac.med.buffalo.edu/  
 VECTOR: pCYPAC2  
 IMPORTANT: This sequence is not the entire insert of clone RP4-579A14. It may be shorter because we sequence overlapping sections only once, except for a 100 base overlap.  
 The true left end of clone RP4-579A14 is at 1 in this sequence. The true left end of clone RP1-230L10 is at 88458 in this sequence. The true right end of clone RP5-826L7 is at 60956 in this sequence.  
 Location/Qualifiers

#### FEATURES

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/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"
/chromosome="6"
/clone_id="RP4-579A14"
/clone_id="RPC1-4"
778. .1087
/note="AluY repeat: matches 1. .305 of consensus"
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1755. .1850
/note="14 copies 4 mer tata 83% conserved"
repeat_region
1853. .1884
/note="16 copies 2 mer at 87% conserved"
repeat_region
1901. .2184
/note="AluSg repeat: matches 1. .284 of consensus"
repeat_region
2380. .2991
/note="36 copies 17 mer 80% conserved"
repeat_region
4022. .4061
/note="MIR repeat: matches 204. .243 of consensus"
repeat_region
4062. .4247
/note="MERSA repeat: matches 1. .189 of consensus"
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4248. .4357
/note="MIR repeat: matches 94. .204 of consensus"
repeat_region
4544. .4676
/note="7 copies 19 mer 88% conserved"
repeat_region
4680. .4831
/note="8 copies 19 mer 89% conserved"
repeat_region
5642. .6058
/note="MSTB repeat: matches 1. .426 of consensus"
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6319. .6326
/note="1318 bases of transposon Tn10 (J01829) removed here. This sequence represents the duplicated flanking sequence of the transposon."
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7142. .7177
/note="9 copies 4 mer acac 88% conserved"
repeat_region
7337. .7538
/note="HAL1 repeat: matches 700. .906 of consensus"
repeat_region
8006. .8248
/note="AluSx repeat: matches 40. .280 of consensus"
repeat_region
8253. .8684
/note="MLT1C repeat: matches 22. .455 of consensus"
repeat_region
8732. .9214
/note="L1PA13 repeat: matches 5663. .6149 of consensus"
repeat_region
10302. .10348
/note="MERS1A repeat: matches 120. .166 of consensus"
repeat_region
10349. .10724
/note="MSTB repeat: matches 1. .426 of consensus"
repeat_region
10732. .11084
/note="MLT1A repeat: matches 1. .365 of consensus"
repeat_region
11085. .11195
/note="MERS1A repeat: matches 1. .114 of consensus"
repeat_region
11489. .11873
/note="MERS7B repeat: matches 1. .403 of consensus"
repeat_region
12045. .12287
/note="AluSx repeat: matches 1. .242 of consensus"
repeat_region
12836. .13137
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repeat_region
13782. .14092
/note="AluY repeat: matches 1. .311 of consensus"
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14290. .14600
/note="AluSx repeat: matches 1. .312 of consensus"
repeat_region
14841. .15132
/note="AluSg repeat: matches 1. .290 of consensus"
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15665. .15962
/note="AluSx repeat: matches 1. .295 of consensus"
repeat_region
16080. .16559
/note="L1MC/D repeat: matches 5319. .5572 of consensus"
repeat_region
16793. .17160
/note="MERS0 repeat: matches 176. .575 of consensus"
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17161. .17285
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17286. .17448
/note="MERS0 repeat: matches 32. .202 of consensus"

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repeat_region
19387. .19574
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19577. .19847
/note="AluSx repeat: matches 1. .281 of consensus"
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19798. .20271
/note="match: GSS: Em:AQ338425"
repeat_region
20076. .20587
/note="MLT1D repeat: matches 1. .498 of consensus"
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21276. .22113
/note="CpG island"
repeat_region
21445. .21527
/evidence=not_experimental
repeat_region
22372. .22666
/note="MLT1A repeat: matches 405. .501 of consensus"
repeat_region
22796. .23027
/note="AluSg repeat: matches 1. .307 of consensus"
repeat_region
23896. .24202
/note="L2 repeat: matches 2425. .2664 of consensus"
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24203. .24563
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24564. .24793
/note="MERSB repeat: matches 1. .362 of consensus"
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26108. .26267
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26109. .26268
/note="40 copies 4 mer tctg 65% conserved"
repeat_region
26281. .26516
/note="80 copies 2 mer gt 65% conserved"
repeat_region
26282. .26563
/note="59 copies 4 mer tctg 71% conserved"
repeat_region
26284. .26517
/note="3 copies 94 mer 73% conserved"
repeat_region
27297. .27445
/note="117 copies 2 mer gt 71% conserved"
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28272. .28443
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29218. .29413
/note="MIR repeat: matches 16. .208 of consensus"
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29407. .29520
/note="MERS8A repeat: matches 1. .191 of consensus"
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29414. .29497
/note="6 copies 19 mer 68% conserved"
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29415. .29504
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repeat_region
29431. .29515
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29522. .29811
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29821. .29870
/note="AluSg repeat: matches 7. .295 of consensus"
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30089. .30632
/note="match: GSS: Em:AQ894713"
repeat_region
32008. .32314
/note="match: GSS: Em:AQ318912"
repeat_region
32331. .32394
/note="AluY repeat: matches 1. .306 of consensus"
repeat_region
32521. .32918
/note="16 copies 4 mer agag 82% conserved"
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33902. .34206
/note="MLT1H repeat: matches 112. .532 of consensus"
repeat_region
34277. .34569
/note="AluY repeat: matches 1. .305 of consensus"
repeat_region
34594. .34840
/note="AluY repeat: matches 7. .297 of consensus"
repeat_region
35170. .35464
/note="MLT1E repeat: matches 308. .568 of consensus"
repeat_region
35465. .35756
/note="MLT1A repeat: matches 65. .365 of consensus"
repeat_region
/note="AluSc repeat: matches 4. .295 of consensus"
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repeat_region 35757..35822
/Note="MT1A1 repeat: matches 1..65 of consensus"
repeat_region 37089..37176
/Note="MER5A repeat: matches 97..187 of consensus"
repeat_region 37728..37832

Query Match 94.4%; Score 17; DB 9; Length 88557;
Best Local Similarity 100.0%; Pred. No. 3;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 GCTGTCCTCCTGTTA 18
|||||
Db 67152 GCTGTCCTCCTGTTA 67136

RESULT 4
AP006146 107979 bp DNA linear PLN 19-MAR-2003
LOCUS Lotus japonicus genomic DNA, chromosome 3, clone:LTJ34M22, TM0282,
DEFINITION complete sequence.
ACCESSION AP006146
VERSION AP006146.1 GI:29122785
KEYWORDS HTG.
SOURCE Lotus japonicus
ORGANISM Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
rosids; eurosids I; Fabales; Fabaceae; Papilionoideae; Loteeae;
Lotus.

REFERENCE
AUTHORS Kaneko, T., Asanizu, E., Kato, T., Sato, S., Nakamura, Y., and Tabata, S.
TITLE Structural Analysis of a Lotus japonicus Genome. III. Sequence
Features and Mapping of Sixty-two TAC Clones Which Cover the 6.7 Mb
Regions of the Genome
JOURNAL DNA Res. 10, 27-33 (2003)
REFERENCE 2 (bases 1 to 107979)
AUTHORS Sato, S.
TITLE Direct Submission
JOURNAL Submitted (12-DEC-2002) Shuei Sato, Kazusa DNA Research Institute,
Department of Plant Gene Research; 2-6-7 Kazusa-kamatari, Kisarazu,
Chiba 252-0818, Japan (E-mail: sato@kazusa.or.jp,
URL: http://www.kazusa.or.jp/, Tel: 81-438-52-3935 (ex. 2337),
Fax: 81-438-52-3934)

FEATURES
source Location/Qualifiers
1..107979
/organism="Lotus japonicus"
/mol_type="genomic DNA"
/db_xref="taxon:34305"
/chromosome="3"
/clone="LTJ34M22"
/clone_lib="JIT library"
/Note="TAC clone: TM0282"
BASE COUNT 33241 a 21293 c 19709 g 33736 t
ORIGIN

Query Match 94.4%; Score 17; DB 8; Length 107979;
Best Local Similarity 100.0%; Pred. No. 3;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 GCTGTCCTCCTGTTA 18
|||||
Db 1793 GCTGTCCTCCTGTTA 1809

RESULT 5
AC128140 226979 bp DNA linear HTG 21-SEP-2002
LOCUS Rattus norvegicus clone CH230-63023, *** SEQUENCING IN PROGRESS
DEFINITION *** 2 unordered pieces.
ACCESSION AC128140
VERSION AC128140.2 GI:23265194
KEYWORDS HTG; HTGS PHASE1; HTGS DRAFT; HTGS_ENRICHED.
SOURCE Rattus norvegicus (Norway rat)

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ORGANISM Rattus norvegicus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
Rattus.

REFERENCE
AUTHORS Muzny, D., Marie, M., Metzger, M., Lee, A., Abramson, S., Adams, C., Alder, J.,
Allen, C., Allen, H., Alsbrooks, S., Amin, A., Anguiano, D.,
Anyalebech, V., Aoyagi, A., Ayodeji, M., Baca, E., Baden, H.,
Baldwin, D., Bandaranaike, D., Barber, M., Barnstead, M., Benahmed, F.,
Biswal, K., Blair, J., Blankenburg, K., Blyth, P., Brown, M.,
Bryant, N., Buhay, C., Burch, P., Burrell, K., Calderon, E.,
Cardenas, V., Carter, K., Cavazos, I., Ceasar, H., Center, A.,
Chacko, J., Chavez, D., Chen, G., Chen, R., Chen, Y., Chen, Z., Chu, J.,
Cleveland, C., Cockrell, R., Cox, C., Coyle, M., Cree, A., D'Souza, L.,
Davila, M., Davis, C., Davy-Carroll, E., De Anda, C., Dederich, D.,
Delgado, O., Denison, S., Deramo, C., Ding, Y., Dinh, H., Divya, K.,
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Hernandez, R., Hines, S., Hladun, S.L., Hodgson, A., Hogues, M.,
Hollins, B., Howells, S., Hulik, S., Hume, J., Idlebird, D., Jackson, A.,
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Liu, J., Liu, W., Liu, Y., London, P., Longacre, S., Lopez, J.,
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Maheshwari, M., Mahindaratne, M., Mahmoud, M., Malloy, K., Mangum, A.,
Mangum, B., Mapua, P., Martin, K., Martin, R., Martinez, E.,
Mawney, S., McLeod, M.P., McNeill, T.Z., Meenen, E.,
Milosavljevic, A., Miner, G., Minja, E., Montemayor, J., Moore, S.,
Morgan, M., Morris, K., Morris, S., Mundasa, M., Murphy, M., Nair, L.,
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Puzo, M., Quiroz, J., Rachlin, E., Reeves, K., Register, M.A., Reigh, R.,
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Rives, C., Rodery, T., Rojas, A., Rose, M., Rose, R., Ritz, S., J.,
Sanders, M., Savery, G., Scherer, S., Scott, G., Shatsman, S., Shen, H.,
Shetty, J., Shvartbeyn, A., Sisson, I., Sitter, C.D., Smajls, D.,
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Steinle, M., Strong, R., Sutton, A., Svatek, A., Tabor, P., Taylor, C.,
Taylor, T., Thomas, N., Thomas, S., Tingey, A., Trejos, Z., Uman, K.,
Valas, R., Vera, V., Villalana, D., Waldron, L., Walker, B., Wang, J.,
Wang, Q., Wang, S., Warren, J., Warren, R., Wei, X., White, F.,
Williams, G., Willison, R., Wlaczky, R., Wooden, H., Worley, K.,
Wright, D., Wright, R., Wu, J., Yakub, S., Yen, J., Yoon, L., Yoon, V.,
Yu, F., Zhang, U., Zhou, J., Zhou, X., Zhao, S., Dunn, D., von
Niederhausern, A., Weiss, R., Smith, D.R., Holt, R.A., Smith, H.O.,
Weinstock, G., and Gibbs, R.A.

TITLE Direct Submission
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 226979)
AUTHORS Worley, K.C.
TITLE Direct Submission
JOURNAL Submitted (19-JUL-2002) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
3 (bases 1 to 226979)

REFERENCE
AUTHORS Rat Genome Sequencing Consortium.
TITLE Direct Submission
JOURNAL Submitted (21-SEP-2002) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
On Sep 21, 2002 this sequence version replaced gi:21908739.
The sequence in this assembly is a combination of BAC based reads
and whole genome shotgun sequencing reads assembled using Atlas
(http://www.hgsc.bcm.tmc.edu/projects/rat/). As a result, the
sequence may extend beyond the ends of the clone and there may be

```

contigs that consist entirely of whole genome shotgun sequence reads. Both end sequences and whole genome shotgun sequence only reads will be indicated in the feature table.

#### ----- Genome Center

Center: Baylor College of Medicine  
Center code: BCM  
Web site: <http://www.hgsc.bcm.tmc.edu/>  
Contact: [hgsc-help@bcm.tmc.edu](mailto:hgsc-help@bcm.tmc.edu)  
Project Information  
Center project name: GYAP  
Center clone name: CH230-63023

#### ----- Summary Statistics

Assembly program: Phrap; version 0.990329  
Consensus quality: 196911 bases at least Q40  
Consensus quality: 199136 bases at least Q30  
Consensus quality: 200537 bases at least Q20  
Estimated insert size: 209329; sum-of-contigs estimation  
Quality coverage: 6x in Q20 bases; sum-of-contigs estimation

-----  
\* NOTE: Estimated insert size may differ from sequence length  
\* (see <http://www.hgsc.bcm.tmc.edu/docs/genbankdraftdata.html>).  
\* NOTE: This is a 'working draft' sequence. It currently  
\* consists of 2 contigs. The true order of the pieces  
\* is not known and their order in this sequence record is  
\* arbitrary. Gaps between the contigs are represented as  
\* runs of N, but the exact sizes of the gaps are unknown.  
\* This record will be updated with the finished sequence  
\* as soon as it is available and the accession number will  
\* be preserved.  
\* 1 224410: contig of 224410 bp in length  
\* 224411 224510: gap of unknown length  
\* 224511 226979: contig of 2469 bp in length.  
Location/Qualifiers

#### FEATURES

##### source

1. 226979

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/mol\_type="genomic DNA"

/db\_xref="taxon:10116"

/clone="CH230-63023"

977..1828

/note="clone boundary"

clone\_end:Sp6

site:ECORI

end\_sequence:BH21166"

85339..86666

/note="wgs contig"

complement[223202..224034]

/note="clone boundary"

clone\_end:T7

site:ECORI

end\_sequence:BH21164"

54404 a 50080 c 47417 g 49761 t 25317 others

BASE COUNT

ORIGIN

Query Match 94.4%; Score 17; DB 2; Length 226979;

Best Local Similarity 100.0%; Pred. No. 2.9;

Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Db 137713 GCTGGTCTACCTGTTA 137697

2 GCTGGTCTACCTGTTA 18

AC096132 231014 bp DNA linear HTG 10-MAY-2003

AC096132 Rattus norvegicus clone CH230-11C18, \*\*\* SEQUENCING IN PROGRESS

AC096132 \*\*\* 2 unordered pieces.

AC096132.6 GI:305222660

HTG; HTGS\_PASSEL; HTGS\_DRAFT; HTGS\_ENRICHED.

Rattus norvegicus (Norway rat)

Rattus norvegicus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

#### REFERENCE

##### AUTHORS

Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;  
Rattus.

1 (bases 1 to 231014)

Muzny, D. Marie, Metzger, M. Lee, Abramson, S., Adams, C., Alder, J.,

Allen, C., Allen, H., Albrooks, S., Aml, A., Anguiano, D.,

Anyadebechi, V., Aoyagi, A., Ayodeji, M., Baca, E., Baden, H.,

Baldwin, D., Bandaranaike, D., Barber, M., Barnstead, M., Benham, F.,

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Bryant, N., Bunay, C., Burch, P., Burrell, K., Calderon, E.,

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Chacko, J., Chavez, D., Chen, G., Chen, R., Chen, Y., Chen, J.,

Cleveland, C., Cockrell, R., Cox, C., Coyle, M., Cre, A., D'Souza, L.,

Davila, M.L., Davis, C., Davy-Carroll, L., De Anda, C., Dederich, D.,

Delgado, O., Denison, S., Deramo, C., Ding, Y., Dinh, H., Divya, K.,

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Fernandez, S., Finley, M., Flagg, N., Forbes, L., Foster, M., Foster, P.,

Fraser, C.M., Gabisi, A., Ganta, R., Garcia, A., Garner, T., Garza, M.,

Gebregiorgis, E., Geer, K., Gill, R., Grady, M., Guerra, M., Guevara, M.,

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Jackson, L., Jacob, L., Jiang, H., Johnson, B., Johnson, R., Jolivet, A.,

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Liu, J., Liu, W., Liu, Y., London, P., Longacre, S., Lopez, J.,

Lorenshew, L., Louised, H., Lozano, R.J., Lu, X., Ma, J.,

Maneshwari, M., Mahindarne, M., Mahmoud, M., Malloy, K., Mangum, A.,

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Mawhney, S., McLeod, M.P., McNeill, T.Z., Meenan, B.,

Milosevic, A., Miner, G., Minja, E., Montemayor, J., Moore, S.,

Morgan, M., Morris, K., Morris, S., Mundasa, M., Murphy, M., Nair, L.,

Nankervis, C., Neal, D., Newton, N., Nguyen, N., Norris, S.,

Nwokedim, O., Okunolu, G., Olamunegbon, A., Pal, S., Parks, K.,

Pasternak, S., Paul, H., Perez, A., Perez, L., Plankoch, C.,

Plopper, F., Polidexter, A., Popovic, D., Primus, E., Pu, L.,

Puazo, M., Quiroz, J., Rachlin, E., Reeves, K., Regier, M.A., Reigh, R.,

Reilly, B., Reilly, M., Ren, Y., Reuter, M., Richards, S., Riggs, F.,

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Sanders, W., Savery, G., Scherer, S., Scott, G., Shatsman, S., Shen, H.,

Shetty, J., Shvartsbeyn, A., Sisson, I., Sitter, C.D., Smajls, D.,

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Valas, R., Vera, V., Villasana, D., Waldron, L., Walker, B., Wang, J.,

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Wright, D., Wright, R., Wu, J., Yakub, S., Yen, U., Yoon, L., Yoon, V.,

Yu, F., Zhang, J., Zhou, J., Zhou, X., Zhao, S., Dunn, D., von

Niederhausen, A., Weiss, R., Smith, D.R., Holt, R.A., Smith, H.O.,

Weinstock, G., and Gibbs, R.A.

Direct Submission

Unpublished

2 (bases 1 to 231014)

Worley, K.C.

Direct Submission

Submitted (17-SEP-2001) Human Genome Sequencing Center, Department

of Molecular and Human Genetics, Baylor College of Medicine, One

Baylor Plaza, Houston, TX 77030, USA

3 (bases 1 to 231014)

Rat Genome Sequencing Consortium.

Submitted (10-MAY-2003) Human Genome Sequencing Center, Department

of Molecular and Human Genetics, Baylor College of Medicine, One

Baylor Plaza, Houston, TX 77030, USA

On May 10, 2003 this sequence version replaced gi:24818180.

The sequence in this assembly is a combination of BAC based reads

and whole genome shotgun sequencing reads assembled using Atlas

(<http://www.hgsc.bcm.tmc.edu/projects/atlas/>). Each contig described

in the feature table below represents a scaffold in the Atlas

assembly (a 'contig-scaffold'). Within each contig-scaffold, and separated

individual sequence contigs are ordered and oriented, and separated

by sized gaps filled with Ns to the estimated size. The sequence may extend beyond the ends of the clone and there may be sequence contigs within a contig-scaffold that consist entirely of whole genome shotgun sequence reads. Both end sequences and whole genome shotgun sequence only contigs will be indicated in the feature table.

#### ----- Genome Center

Center: Baylor College of Medicine  
Center code: BCM  
Web site: <http://www.hgsc.bcm.tmc.edu/>  
Contact: [hgsc-help@bcm.tmc.edu](mailto:hgsc-help@bcm.tmc.edu)

#### ----- Project Information

Center project name: GEMO  
Center clone name: CH230-11C18

#### ----- Summary Statistics

Assembly program: Atlas 3.0  
Consensus quality: 208633 bases at least Q40  
Consensus quality: 212147 bases at least Q30  
Consensus quality: 214386 bases at least Q20  
Estimated insert size: 214070; sum-of-contigs estimation  
Quality coverage: 6x in Q20 bases; sum-of-contigs estimation

\* NOTE: Estimated insert size may differ from sequence length  
\* (see [http://www.hgsc.bcm.tmc.edu/docs/Genbank\\_draft\\_data.html](http://www.hgsc.bcm.tmc.edu/docs/Genbank_draft_data.html)).  
\* NOTE: This is a 'working draft' sequence. It currently  
\* consists of 2 contigs. The true order of the pieces  
\* is not known and their order in this sequence record is  
\* arbitrary. Gaps between the contigs are represented as  
\* runs of N, but the exact sizes of the gaps are unknown.  
\* This record will be updated with the finished sequence  
\* as soon as it is available and the accession number will  
\* be preserved.

#### FEATURES

##### source

1 229193: contig of 229193 bp in length  
\* 229194 229293: gap of unknown length  
\* 229294 231014: contig of 1721 bp in length.  
Location/Qualifiers

1. 231014

/organism="Rattus norvegicus"

/db\_xref="taxon:10116"

/clone="CH230-11C18"

1. 1031

/note="wgs end extension"

/clone\_end:"T7"

2512..3242

/note="clone\_boundary"

/clone\_end:"T7"

site:ECORI

end sequence: BH320811"

23368..24820

/note="wgs contig"

121675..123616

/note="wgs contig"

124270..126190

/note="wgs contig"

169881..171027

/note="wgs contig"

complement(228041..228684)

/note="clone\_boundary"

clone\_end:Sp6

site:ECORI

end sequence: BH320830"

61558 a 48903 c 47136 g 58505 t 14912 others

BASE COUNT

ORIGIN

Query Match

Best Local Similarity 94.4%; Score 17; DB 2; Length 231014;

Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

1 GCGTGGTGTACCTGTT 17

DB 122048 GCGTGGTGTACCTGTT 122064

#### RESULT 7

AC140198

LOCUS

DEFINITION

AC140198

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

REFERENCE

AUTHORS

TITLE

JOURNAL

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

AC140198 238552 bp DNA linear HTG 01-MAR-2003  
Mus musculus chromosome UNK clone RP23-255C4, WORKING DRAFT  
SEQUENCE, 4 unordered pieces.

AC140198.2 GI:28626914  
HTG; HTGS PHASE1; HTGS DRAFT; HTGS\_FULLTOP.

Mus musculus (house mouse)

Mus musculus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

1 (bases 1 to 238552)

McPherson,J.D. and Waterston,R.H.

The sequence of Mus musculus clone

Unpublished

2 (bases 1 to 238552)

McPherson,J.D. and Waterston,R.H.

Direct Submissions

Submitted (23-FEB-2003) Genome Sequencing Center, 4444 Forest Park

Parkway, St. Louis, MO 63108, USA

3 (bases 1 to 238552)

McPherson,J.D. and Waterston,R.H.

Direct Submissions

Submitted (01-MAR-2003) Genome Sequencing Center, 4444 Forest Park

Parkway, St. Louis, MO 63108, USA

On Mar 1, 2003 this sequence version replaced gi:28475433.

----- Genome Center -----  
Center: Washington University Genome Sequencing Center  
Center code: WUGSC  
Web site: <http://genome.wustl.edu/gsc/index.shtml>  
Contact: [submissions@wustl.edu](mailto:submissions@wustl.edu)

----- Project Information -----  
Center project name: M\_BA0255C04

----- Summary Statistics -----

Sequencing vector: MJ3; 0%

Chemistry: Dye-terminator Big Dye; 100% of reads

Assembly program: Phrap; version 0.990319

Consensus quality: 236489 bases at least Q40

Consensus quality: 236878 bases at least Q30

Consensus quality: 237189 bases at least Q20

Insert size: 183000; agarose-fp

Insert size: 248898; sum-of-contigs

Quality coverage: 16.05 in Q20 bases; agarose-fp

Quality coverage: 12.12 in Q20 bases; sum-of-contigs

\* NOTE: This is a 'working draft' sequence. It currently  
\* consists of 4 contigs. The true order of the pieces  
\* is not known and their order in this sequence record is  
\* arbitrary. Gaps between the contigs are represented as  
\* runs of N, but the exact sizes of the gaps are unknown.  
\* This record will be updated with the finished sequence  
\* as soon as it is available and the accession number will  
\* be preserved.

1 2229: contig of 2229 bp in length

2230 2339: gap of unknown length

2230 28154: contig of 25825 bp in length

28155 28254: gap of unknown length

28255 98428: contig of 70174 bp in length

98429 98528: gap of unknown length

98529 238552: contig of 140024 bp in length.

#### FEATURES

##### source

1. 238552

/organism="Mus musculus"

/mol\_type="genomic DNA"

/db\_xref="taxon:10090"

/chromosome="UNK"

/clone="RP23-255C4"



```

misc_feature      1. 2229
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misc_feature      2330. 28154
                  /note="assembly_name:Contig24"
misc_feature      28255. 98428
                  /note="assembly_name:Contig25"
misc_feature      98529. 238552
                  /note="assembly_name:Contig26"
BASE COUNT      69322 a 53309 c 49714 g 65901 t      306 others
ORIGIN
Query Match      94.4%; Score 17; DB 2; Length 238552;
Best Local Similarity 100.0%; Pred. No. 2.9;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy      2 GCTGTCACCTGTTA 18
Db      155734 GCTGTCACCTGTTA 155750

RESULT 8
AC131614
LOCUS      AC131614      244424 bp      DNA      linear      HTG 20-NOV-2002
DEFINITION      Rattus norvegicus clone CH230-284C9, WORKING DRAFT SEQUENCE, 3
unordered pieces.
AC131614
AC131614.3 GI:25138981
HTG: HTGS PHASE1: HTGS DRAFT: HTGS FULLTOP.
Rattus norvegicus (Norway rat)
Rattus norvegicus
Eukaryote; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
Rattus.
1 (bases 1 to 244424)
Mizny,D.Marie, Metzker,M.Lee, Abramzon,S., Adams,C., Alder,J.,
Allen,C., Allen,H., Alsbrooks,S., Amin,A., Angiano,D.,
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Fraser,C.M., Gabisti,A., Ganta,R., Garcia,A., Garner,T., Garza,M.,
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Hollins,B., Howells,S., Huliy,S., Hume,J., Idlebird,D., Jackson,A.,
Jackson,B., Jacob,L., Jiang,H., Johnson,B., Johnson,R., Jollivet,A.,
Karpathy,S., Kelly,S., Kelly,S., Khan,Z., King,L., Kovar,C.,
Kowis,C., Kraft,C.L., Lebow,H., Levan,J., Lewis,L., Li,Z., Liu,J.,
Liu,J., Liu,W., Liu,Y., London,P., Longacre,S., Lopez,J.,
Lorensunewa,L., Louieged,H., Lozada,R.J., Lu,X., Ma,J.,
Maheshwari,M., Mahindaratne,M., Mahmoud,M., Malloy,K., Mangum,A.,
Mangum,B., Mapua,P., Martin,K., Martin,R., Martinez,E.,
Mawhney,S., McLeod,M.P., McNeill,T.Z., Meenen,E.,
Milogajlevic,A., Miner,G., Minja,E., Montemayor,J., Moore,S.,
Morgan,M., Morris,K., Morris,S., Munidasa,M., Murphy,M., Nair,L.,
Nankervis,C., Neal,D., Newton,N., Nguyen,N., Norris,S.,
Nwaokwelenh,O., Okwuonu,G., Olarinpoosoon,A., Pal,S., Parks,K.,
Pasternak,S., Paul,H., Perez,A., Perez,L., Plamnick,C.,
Plopper,F., Poindexter,A., Popovic,D., Primus,E., Pu,L.-L.,
Puzio,B., Quiroz,J., Rachlin,E., Reeves,K., Regier,M.A., Reigh,R.,
Reilly,B., Reilly,M., Ren,Y., Reuter,M., Richards,S., Riggs,F.,
Rives,C., Rodkey,T., Rojas,A., Rose,M., Rose,R., Ruiz,S.J.,
Sanders,M., Savary,G., Scherer,S., Scott,G., Shatsman,S., Shen,H.,
Shetty,D., Shvartsbeyn,A., Sisson,I., Sitter,C.D., Smajs,D.,

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Sneed,A., Sodergren,E., Song,X.-Z., Sorelle,R., Sosa,J.,
Steinle,M., Strong,R., Sutton,A., Svatek,A., Tabor,P., Taylor,C.,
Taylor,T., Thomas,N., Thomas,S., Tingey,A., Trejos,Z., Umanai,K.,
Valas,R., Vera,V., Villaseana,D., Waldron,L., Walker,B., Wang,J.,
Wang,Q., Wang,S., Warren,J., Warren,R., Wei,X., White,F.,
Williams,G., Wilson,R., Wleczyk,R., Wodden,H., Wortley,K.,
Wright,D., Wright,R., Wu,J., Yakub,S., Yen,J., Yoon,L., Yoon,V.,
Yu,F., Zhang,J., Zhou,J., Zhou,X., Zhou,S., Dunn,D., von
Niederhausen,A., Weiss,R., Smith,D.R., Holt,R.A., Smith,H.O.,
Weinstock,G. and Gibbs,R.A.
Direct Submission
Unpublished
2 (bases 1 to 244424)
REFERENCE
Rat Genome Sequencing Consortium.
AUTHORS
Rat Genome Sequencing Consortium.
TITLE
Submitted (25-NOV-2002) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
3 (bases 1 to 244424)
REFERENCE
Rat Genome Sequencing Consortium.
AUTHORS
Rat Genome Sequencing Consortium.
TITLE
Submitted (20-NOV-2002) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
COMMENT
The sequence in this assembly is a combination of BAC based reads
and whole genome shotgun sequencing reads assembled using Atlas
(http://www.hgsc.bcm.tmc.edu/projects/rat/). Each contig described
in the feature table below represents a scaffold in the Atlas
assembly (a 'contig-scaffold'). Within each contig-scaffold,
individual sequence contigs are ordered and oriented, and separated
by sized gaps filled with Ns to the estimated size. The sequence
may extend beyond the ends of the clone and there may be sequence
contigs within a contig-scaffold that consist entirely of whole
genome shotgun sequence reads. Both end sequences and whole genome
shotgun sequence only contigs will be indicated in the feature
table.
----- Genome Center
Center: Baylor College of Medicine
Center code: BCM
Web site: http://www.hgsc.bcm.tmc.edu/
Contact: hgsc-help@bcm.tmc.edu
----- Project Information
Center project name: KCCL
Center clone name: CH230-284C9
----- Summary Statistics
Assembly program: Phrap, version 0.990329
Consensus quality: 22132 bases at least Q40
Consensus quality: 22413 bases at least Q30
Consensus quality: 225930 bases at least Q20
Estimated insert size: 226559; sum-of-contigs estimation
Quality coverage: 7x in Q20 bases; sum-of-contigs estimation
* NOTE: Estimated insert size may differ from sequence length
* (see http://www.hgsc.bcm.tmc.edu/docs/Genbank_draft_data.html).
* NOTE: This is a 'working draft' sequence. It currently
* consists of 3 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
* 1 235352: contig of 235352 bp in length
* 235353 235452: gap of unknown length
* 235453 240268: contig of 4816 bp in length
* 240269 240368: gap of unknown length
* 240369 244424: contig of 4056 bp in length.
FEATURES
source
1. 244424
/organism="Rattus norvegicus"
/mol_type="genomic DNA"
/db_xref="taxon:10116"

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misc_feature      /clone="CH230-284C9"
                  1..1120
                  /note="wgs_end_extension
clone_end:T7"
misc_feature      1249..2544
                  /note="wgs_end_extension
clone_end:T7"
misc_feature      complement(3470..4259)
                  /note="clone_boundary
clone_end:T7
                  site:
                  end_sequence:RXAHE177GB"
BASE COUNT      56117 a 54386 c 56712 g 59935 t 17274 others
ORIGIN
Query Match      94.4% Score 17; DB 2; Length 244424;
Best Local Similarity 100.0%; Pred. No. 2.9;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      2 GCGGAGTGCACCTGTTA 18
        |||||
Db      203097 GCTGGTGCACCTGTTA 203113

RESULT 9
AC123215/c
LOCUS      AC123215
DEFINITION Rattus norvegicus clone CH230-118B7, *** SEQUENCING IN PROGRESS
***
AC123215      244636 bp DNA linear HTG 21-SEP-2002
Rattus norvegicus clone CH230-118B7, *** SEQUENCING IN PROGRESS
AC123215      GI:23265875
HTG: HTGS PHASE2; HTGS_DRAFT; HTGS_ENRICHED.
SOURCE      Rattus norvegicus (Norway rat)
ORGANISM      Rattus norvegicus
              Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
              Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
              Rattus.
              1 (bases 1 to 244636)
              Muzny,D,Marie, Metzker,M,Lee, Abramson,S, Adams,C, Alder,J,
              Allen,C, Allen,H, Alsbrooks,S, Amin,A, Argulano,D,
              Anyalebechi,V, Aoyagi,A, Ayodeji,M, Baca,E, Baden,H,
              Baldwin,D, Bandaranaike,D, Barber,M, Barnstead,M, Benahmed,F,
              Biewlo,K, Blair,J, Blankenburg,K, Blyth,P, Brown,M,
              Bryant,N, Buhay,C, Burch,P, Burrell,K, Calderon,E,
              Cardenas,V, Carter,K, Cavazos,I, Caesar,H, Center,A,
              Chacko,J, Chavez,D, Chen,G, Chen,R, Chen,Y, Chen,Z, Chu,J,
              Cleveland,C, Cockrell,R, Cox,C, Coyle,M, Cree,A, D'Souza,L,
              Davila,M,L, Davis,C, Davy-Carroll,L, De Anda,C, Dedereich,D,
              Delgado,O, Denson,S, Deramo,C, Ding,Y, Dinh,H, Divya,K,
              Draper,H, Dugan-Rocha,S, Dunn,A, Durbin,K, Duval,B, Eaves,K,
              Egan,A, Escotto,M, Eugene,C, Evans,C,A, Falls,T, Fan,G,
              Fernandez,S, Finley,M, Flegg,N, Forbes,L, Foster,M, Foster,P,
              Frazer,C,M, Gabisi,A, Ganta,R, Garcia,A, Garner,T, Garza,M,
              Gebregeorgis,E, Geer,K, Gill,R, Grady,M, Guerra,M, Guevara,W,
              Gunaratne,P, Haaland,W, Hamill,C, Hamilton,C, Hamilton,K,
              Harvey,X, Havlak,P, Hawes,A, Henderson,N, Hernandez,J,
              Hernandez,R, Hines,S, Hladun,S,L, Hodgson,A, Hogue,M,
              Hollins,B, Howells,S, Huily,S, Hume,J, Idlebird,D, Jackson,A,
              Jackson,B, Jacob,L, Jiang,H, Johnson,B, Johnson,R, Jolyet,A,
              Karpathy,S, Kelly,S, Kelly,S, Khan,Z, King,L, Kovar,C,
              Kowls,C, Kraft,C,L, Lebow,H, Levan,Z, Lewis,L, Li,Z, Liu,J,
              Liu,J, Liu,W, Liu,Y, London,P, Longacre,S, Lopez,J,
              Lorensuewa,L, Louieged,H, Lozano,R,J, Lu,X, Ma,J,
              Maheshwari,M, Mahindaratne,M, Mahmoud,M, Malloy,K, Mangum,A,
              Mangum,B, Mapa,P, Martin,K, Martin,R, Martinez,E,
              Mawhood,S, McLeod,M,P, McNeill,T,Z, Meenan,E,
              Mlisaajevic,A, Miner,G, Minja,E, Montemayor,J, Moore,S,
              Morgan,M, Morris,K, Morris,S, Munidesa,M, Murphy,M, Nair,L,
              Nankervis,C, Neal,D, Newton,N, Nguyen,N, Norris,S,
              Nwaokemelehen,O, Okwuonu,G, Olarunpusagoon,A, Pal,S, Parks,K,
              Pasternak,S, Paul,H, Perez,A, Perez,L, Pfannkuch,C,
              Plodper,F, Poindexter,A, Popovic,D, Primus,E, Pu,L,-L,
              Puzo,M, Quiroz,J, Rachlin,E, Reeves,K, Regier,M,A, Reigh,R,

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TITLE
JOURNAL
REFERENCE
AUTHORS
TITLE
JOURNAL
REFERENCE
AUTHORS
TITLE
JOURNAL
COMMENT
Center: Baylor College of Medicine
Center code: BCM
Web site: http://www.hgsc.bcm.tmc.edu/
Contact: hgsc-help@bcm.tmc.edu
Project Information
Center project name: GSCP
Center clone name: CH230-118B7
----- Summary Statistics -----
Assembly program: Phrap version 0.990329
Consensus quality: 223663 bases at least Q40
Consensus quality: 225326 bases at least Q30
Consensus quality: 226426 bases at least Q20
Estimated insert size: 241958; sum-of-coverage estimation
Quality coverage: 4x in Q20 bases; sum-of-coverage estimation
----- NOTE -----
* NOTE: Estimated insert size may differ from sequence length
  (see http://www.hgsc.bcm.tmc.edu/docs/Genbank_draft_data.html).
* NOTE: This is a 'working draft' sequence. It currently
  * consists of 1 contigs. Gaps between the contigs
  * are represented as runs of N. The order of the pieces
  * is believed to be correct as given, however the sizes
  * of the gaps between them are based on estimates that have
  * been provided by the submitter.
  * This sequence will be replaced
  * by the finished sequence as soon as it is available and
  * the accession number will be preserved.
  1 244636: contig of 244636 bp in length.
Location/Qualifiers
1..244636
/organism="Rattus norvegicus"
/mol_type="genomic DNA"
/db_xref="taxon:10116"
/clone="CH230-118B7"
1034..1835
/note="clone boundary
clone_end:Sp6

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misc_feature      site:ECORI
                  end sequence:BM285202"
                  complement(244009..244636)
                  /note="Clone_boundary
                  clone_end:17
                  site:ECORI
                  end sequence:BM285200"
BASE COUNT      62840 a 49128 c 49531 g 65556 t 17581 others
ORIGIN
Query Match      94.4%; Score 17; DB 2; Length 244636;
Best Local Similarity 100.0%; Pred. No. 2.9;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

1 GGCTGTGTCTACCTGTT 17
Db 45298 GGCTGTGTCTACCTGTT 45282

RESULT 10
AC103080/c
LOCUS      AC103080      248793 bp      DNA      linear      HTG 13-MAY-2003
DEFINITION Rattus norvegicus clone CH230-22501, WORKING DRAFT SEQUENCE, 5
unordered pieces.
AC103080
AC103080.6 GI:30579747
HTG: HTGS PHASE1: HTGS DRAFT: HTGS_FULLTOP.
Rattus norvegicus (Norway rat)
Rattus norvegicus
Eukaryote; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
Rattus.
1 (bases 1 to 248793)
Muzny,D,Marle,M,Metzker,M,Lee,A,Abramson,S,Adams,C,Alder,J,
Allen,C,Allen,H,Aisbrooks,S,Amin,A,Anguiano,D,
Ayala-Becchi,V,Aoyagi,A,Ayodeji,M,Baca,E,Baden,H,
Baldwin,D,Bandaranaike,D,Barber,M,Barnstead,M,Benahmed,F,
Biswal,K,Blair,J,Blankenburg,K,Blyth,P,Brown,M,
Bryan,N,Buhay,C,Burch,P,Burrell,K,Calderson,E,
Cardenas,V,Carter,K,Cavazos,I,Ceasar,H,Center,A,
Chacko,J,Chavez,D,Chen,G,Chen,R,Chen,Y,Chen,Z,Chu,J,
Cleveland,C,Cockrell,R,Cox,C,Coyle,M,Cree,A,D'Souza,L,
Davila,M,L,Davis,C,Davy-Carroll,L,De Anda,C,Decker,D,
Delgado,O,Denson,S,Deramo,C,Ding,Y,Dinh,H,Ditaya,K,
Draper,H,Dugan-Rocha,S,Dunn,A,Durbin,K,Duval,B,Eaves,K,
Egan,A,Escotto,M,Eugene,C,Evans,C,A,Falls,T,Fan,G,
Fernandez,S,Finley,M,Flagg,N,Fordes,L,Foster,M,Foster,P,
Fraser,C,M,Gabisi,A,Ganta,R,Garcia,A,Garnier,T,Garza,M,
Gibbs,G,Gilbert,J,Gier,K,Gill,R,Grady,M,Guerra,W,Guevara,W,
Gunaratne,P,Halland,W,Hamil,C,Hamilton,C,Hamilton,K,
Harvey,Y,Havlak,P,Hawes,A,Henderson,N,Hernandez,J,
Hollander,R,Hines,S,Hladun,S,L,Hodgson,A,Hogues,M,
Hollins,B,Howell,S,Huylk,S,Hume,J,Idelbird,D,Jackson,A,
Jackson,L,Jacob,L,Jiang,H,Johnson,B,Johnson,R,Jolivet,A,
Karpathy,S,Kelly,S,Kelly,S,Khan,Z,King,L,Kovar,C,
Kowis,C,Kraft,C,L,Lebow,H,Levan,J,Lewis,L,Li,Z,Liu,J,
Liu,J,Liu,W,Liu,Y,London,P,Longacre,S,Lopez,J,
Lorenzini,L,Louisege,H,Lorado,R,J,Lu,X,Ma,J,
Maheshwari,M,Mahindratne,M,Mahmoud,M,Malloy,K,Mangum,A,
Mangum,B,Mapua,P,Martin,K,Martin,R,Martinez,E,
Mawhney,S,McLeod,M,P,McNeill,T,Z,Meenen,E,
Mlosoajlevic,A,Miner,G,Ming,M,Munidas,M,Murphy,J,Moore,S,
Morgan,M,Morris,K,Morris,S,Munidas,M,Murphy,J,Moore,S,
Nankervis,C,Neal,D,Newton,N,Nguyen,N,Norris,S,
Nwaokoleh,O,Okinonu,G,Olatunbosun,A,Pal,S,Parker,K,
Pasternak,S,Paul,H,Perez,A,Perez,L,Pfannkuch,C,
Plappier,F,Polindexter,A,Popovic,D,Primus,E,Pu,L-L,
Puzos,M,Quiroz,J,Rachlin,E,Reeves,K,Regier,M,A,Reigh,R,
Riley,B,Riley,M,Ren,Y,Reuter,M,Richard,S,Riggs,F,
Rivers,C,Rodkey,T,Rojas,A,Rose,M,Rose,R,Ruiz,S,J,
Sanders,M,Savery,G,Scherrer,S,Scott,G,Shamsan,S,Shen,H,
Shetty,J,Shvartsbeyn,A,Sisson,I,Sitter,C,D,Smajs,D,
Sneed,A,Sodergren,E,Song,X-Z,Sorelle,R,Sosa,J,

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TITLE
JOURNAL
AUTHORS
REFERENCE
JOURNAL
TITLE
Submitted (24-NOV-2001) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
3 (bases 1 to 248793)
Rat Genome Sequencing Consortium.
Direct Submission
Submitted (13-MAY-2003) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
On May 13, 2003 this sequence version replaced gi:23123516.
The sequence in this assembly is a combination of BAC based reads
and whole genome shotgun sequencing reads assembled using Atlas
(http://www.hgsc.bcm.tmc.edu/projects/rat/). Each contig described
in the feature table below represents a scaffold in the Atlas
assembly (a 'contig-scaffold'). Within each contig-scaffold,
individual sequence contigs are ordered and oriented, and separated
by sized gaps filled with Ns to the estimated size. The sequence
may extend beyond the ends of the clone and there may be sequence
contigs within a contig-scaffold that consist entirely of whole
genome shotgun sequence reads. Both end sequences and whole genome
shotgun sequence only contigs will be indicated in the feature
table.
----- Genome Center
Center: Baylor College of Medicine
Center code: BCM
Web site: http://www.hgsc.bcm.tmc.edu/
Contact: hgsc-help@bcm.tmc.edu
----- Project Information
Center project name: GJBC
Center clone name: CH230-22501
----- Summary Statistics
Assembly program: Atlas 3.0:
Consensus quality: 213558 bases at least Q40
Consensus quality: 215685 bases at least Q30
Consensus quality: 217422 bases at least Q20
Estimated insert size: 218947; sum-of-contigs estimation
Quality coverage: 8x in Q20 bases; sum-of-contigs estimation
-----
* NOTE: Estimated insert size may differ from sequence length
(see http://www.hgsc.bcm.tmc.edu/docs/genbank_draft_data.html).
* NOTE: This is a 'working draft' sequence. It currently
consists of 5 contigs. The true order of the pieces
is not known and their order in this sequence record is
arbitrary. Gaps between the contigs are represented as
runs of N, but the exact sizes of the gaps are unknown.
This record will be updated with the finished sequence
as soon as it is available and the accession number will
be preserved.
1 243312: contig of 243312 bp in length
243313 243412: gap of unknown length
243413 244623: contig of 1211 bp in length
244624 244723: gap of unknown length
244724 245933: contig of 1210 bp in length
245934 246033: gap of unknown length
246034 247253: contig of 1220 bp in length
247254 247353: gap of unknown length
247354 248793: contig of 1440 bp in length.
1.248793
Location/Qualifiers

```

FEATURES  
source

```

/misc_feature
/misc_feature
misc_feature
BASE COUNT      59791 a 53488 c 51802 g 54201 t 29511 others
ORIGIN
Query Match
Best Local Similarity      94.4%; Score 17; DB 2; Length 248793;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY      2 GCTGCTGCACCTGTTA 18
|||||
Db      183996 GCTGCTGCACCTGTTA 183980
9

RESULT 11
LOCUS      126102
DEFINITION Sequence_28 from patent US 556772.
ACCESSION  126102
VERSION    126102.1 GI:1605972
KEYWORDS
SOURCE     Unknown.
ORGANISM   Unknown.
REFERENCE   1 (bases 1 to 20)
AUTHORS    Sorge,J.A. and Mullinax,R.L.
TITLE      Polymerase-compositions-and-uses-thereof
JOURNAL    Patent: US 556772-A 28 17-SEP-1996;
FEATURES
source     1..20
              /organism="unknown"
BASE COUNT      2 a      4 c      7 g      7 t
ORIGIN
Query Match
Best Local Similarity      88.9%; Score 16; DB 6; Length 20;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY      1 GGCTGGTGTCACTGT 16
|||||
Db      1 GGCTGGTGTCACTGT 16

RESULT 12
LOCUS      AR032934/c
DEFINITION AR032934
ACCESSION  AR032934
VERSION    AR032934.1 GI:5948539
KEYWORDS
SOURCE     Unknown.
ORGANISM   Unknown.
REFERENCE   1 (bases 1 to 50)
AUTHORS    Edwards,C.A., Cantor,C.R., Andrews,B.M., Turin,L.M. and Fry,K.E.
TITLE      Method of determining DNA sequence preference of a DNA-binding molecule.
JOURNAL    Patent: US 5869241-A 546 09-FEB-1999;
FEATURES
source     1..50
              /organism="unknown"

```

ORIGIN	BASE COUNT	18 a	14 c	10 g	8 t
Query Match	88.9%; Score 16; DB 6; Length 50;				
Best Local Similarity	100.0%; Pred. No. 16;				
Matches	16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;				
QY	1 GGCTGGTGTACCTGT 16				
DB	41 GGCTGGTGTACCTGT 26				
RESULT 13					
LOCUS	AR209598	50 bp			
DEFINITION	Sequence 546 from patent US 6384208.				
ACCESSION	AR209598				
VERSION	AR209598.1				
KEYWORDS	GI:21511065				
SOURCE	Unknown.				
ORGANISM	Unclassified.				
REFERENCE	1 (bases 1 to 50)				
AUTHORS	Edwards,C.A., Cantor,C.R., Andrews,B.M., Turin,L.M. and Fry,K.E.				
TITLE	Sequence directed DNA binding molecules compositions and methods				
JOURNAL	Patent: US 6384208-A 546 07-MAY-2002;				
FEATURES	Location/Qualifiers				
source	1..50				
	/organism="unknown"				
BASE COUNT	18 a	14 c	10 g	8 t	
ORIGIN					
Query Match	88.9%; Score 16; DB 6; Length 50;				
Best Local Similarity	100.0%; Pred. No. 16;				
Matches	16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;				
QY	1 GGCTGGTGTACCTGT 16				
DB	41 GGCTGGTGTACCTGT 26				
RESULT 14					
LOCUS	129674	50 bp			
DEFINITION	Sequence 546 from patent US 5578444.				
ACCESSION	129674				
VERSION	129674.1				
KEYWORDS	GI:1820465				
SOURCE	Unknown.				
ORGANISM	Unclassified.				
REFERENCE	1 (bases 1 to 50)				
AUTHORS	Edwards,C.A., Cantor,C.R., Andrews,B.M., Turin,L.M. and Fry,K.E.				
TITLE	Sequence-directed DNA-binding molecules compositions and methods				
JOURNAL	Patent: US 5578444-A 546 26-NOV-1996;				
FEATURES	Location/Qualifiers				
source	1..50				
	/organism="unknown"				
BASE COUNT	18 a	14 c	10 g	8 t	
ORIGIN					
Query Match	88.9%; Score 16; DB 6; Length 50;				
Best Local Similarity	100.0%; Pred. No. 16;				
Matches	16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;				
QY	1 GGCTGGTGTACCTGT 16				
DB	41 GGCTGGTGTACCTGT 26				
RESULT 15					
LOCUS	191348	50 bp			
DEFINITION	Sequence 546 from patent US 5578444.				
ACCESSION	191348				
VERSION	191348.1				
KEYWORDS	GI:1820465				
SOURCE	Unknown.				
ORGANISM	Unclassified.				
REFERENCE	1 (bases 1 to 50)				
AUTHORS	Edwards,C.A., Cantor,C.R., Andrews,B.M., Turin,L.M. and Fry,K.E.				
TITLE	Sequence-directed DNA-binding molecules compositions and methods				
JOURNAL	Patent: US 5578444-A 546 26-NOV-1996;				
FEATURES	Location/Qualifiers				
source	1..50				
	/organism="unknown"				
BASE COUNT	18 a	14 c	10 g	8 t	
ORIGIN					
Query Match	88.9%; Score 16; DB 6; Length 50;				
Best Local Similarity	100.0%; Pred. No. 16;				
Matches	16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;				
QY	1 GGCTGGTGTACCTGT 16				
DB	41 GGCTGGTGTACCTGT 26				
RESULT 16					
LOCUS	191348	50 bp			
DEFINITION	Sequence 546 from patent US 5578444.				
ACCESSION	191348				
VERSION	191348.1				
KEYWORDS	GI:1820465				
SOURCE	Unknown.				
ORGANISM	Unclassified.				

DEFINITION Sequence 546 from patent US 5726014.  
ACCESSION 191348  
VERSION 191348.1 GI:3935818  
KEYWORDS  
SOURCE Unknown.  
ORGANISM Unknown.  
REFERENCE 1 (bases 1 to 50)  
AUTHORS Edwards, C.A., Cantor, C.R., Andrews, B.M. and Turin, L.M.  
TITLE Screening assay for the detection of DNA-binding molecules  
JOURNAL Patent: US 5726014-A 546 10-MAR-1998;  
FEATURES  
source 1.50  
BASE COUNT 18 a 14 c 10 g 8 t  
ORIGIN  
Query Match 88.9%; Score 16; DB 6; Length 50;  
Best Local Similarity 100.0%; Pred. No. 16;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 1 GGCTGGTGCACCTGT 16  
Db 41 GGCTGGTGCACCTGT 26  
RESULT 16  
LOCUS BD025521/c 435 bp DNA linear PAT 27-AUG-2002  
DEFINITION Sequence tag and encoded human protein.  
ACCESSION BD025521  
VERSION BD025521.1 GI:22566744  
KEYWORDS JP 2001269182-A/1767.  
SOURCE Homo sapiens (human)  
ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.  
REFERENCE 1 (bases 1 to 435)  
AUTHORS Edwards, J.B.D.M., Duclair, E. and Jordan, J.Y.  
TITLE Sequence tag and encoded human protein.  
JOURNAL Patent: JP 2001269182-A 1767 02-OCT-2001;  
COMMENT GENSET  
OS Homo sapiens (human)  
PN JP 2001269182-A/1767  
PD 02-OCT-2001  
PR 24-FEB-2000 JP 2000118773  
PI 26-FEB-1999 US 60/122487  
PT JEAN BAPTISTE DOMAS MILNE EDWARDS, EIMERIC DUCLAIR, JEAN YVES  
PC C12N15/09, C07K14/435, C07K16/18, C12N1/15, C12N1/19, C12N1/21, PC  
C12N5/10,  
PC C12P21/02, C12P21/08, C12Q1/68//G06F17/30, C12N15/00, C12N5/00, PC  
G06F15/40  
CC  
FH Key Location/Qualifiers  
FT CDS 60..434..  
FEATURES  
source 1..435 Location/Qualifiers  
BASE COUNT 105 a 114 c 133 g 79 t 4 others  
ORIGIN  
Query Match 88.9%; Score 16; DB 6; Length 435;  
Best Local Similarity 100.0%; Pred. No. 15;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 1 GGCTGGTGCACCTGT 16  
Db 186 GGCTGGTGCACCTGT 171

RESULT 17  
LOCUS AX321046/c 455 bp DNA linear PAT 15-DEC-2001  
DEFINITION Sequence 63 from Patent WO0177168.  
ACCESSION AX321046  
VERSION AX321046.1 GI:17904325  
KEYWORDS  
SOURCE Homo sapiens (human)  
ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.  
REFERENCE 1 (bases 1 to 963)  
AUTHORS Lodes, M.J., Wang, T., Mohamath, R. and Indirias, C.Y.  
TITLE Compositions and methods for the therapy and diagnosis of lung cancer  
JOURNAL Patent: WO 0177168-A 63 18-OCT-2001;  
CORIXA CORPORATION (US)  
FEATURES  
source 1..455 Location/Qualifiers  
BASE COUNT 122 a 116 c 138 g 78 t 1 others  
ORIGIN  
Query Match 88.9%; Score 16; DB 6; Length 455;  
Best Local Similarity 100.0%; Pred. No. 15;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 1 GGCTGGTGCACCTGT 16  
Db 177 GGCTGGTGCACCTGT 162  
RESULT 18  
LOCUS BT006889/c 963 bp mRNA linear PRI 13-MAY-2003  
DEFINITION Homo sapiens eukaryotic translation initiation factor 3, subunit 4  
ACCESSION BT006889  
VERSION BT006889.1 GI:30582616  
KEYWORDS FLI\_CDN.  
SOURCE Homo sapiens (human)  
ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.  
REFERENCE 1 (bases 1 to 963)  
AUTHORS Kainline, N., Chen, X., Rolfs, A., Halleck, A., Hines, L., Eisenstein, S.,  
Koundinya, M., Raphael, J., Moreira, D., Kelley, T., Labaer, J., Lin, Y.,  
Phelan, M. and Farmer, A.  
TITLE Cloning of human full-length CDS in BD Creator(TM) System Donor  
vector  
JOURNAL Unpublished  
REFERENCE 2 (bases 1 to 963)  
AUTHORS Kainline, N., Chen, X., Rolfs, A., Halleck, A., Hines, L., Eisenstein, S.,  
Koundinya, M., Raphael, J., Moreira, D., Kelley, T., Labaer, J., Lin, Y.,  
Phelan, M. and Farmer, A.  
TITLE Direct Submission  
JOURNAL Submitted (13-MAY-2003) BD Biosciences Clontech, 1020 East Meadow  
Circle, Palo Alto, CA 94303, USA  
COMMENT This CDS clone is a part of a collection of human full length  
expression clones generated by BD Biosciences Clontech and the  
Harvard Institute of Proteomics. Each CDS has been cloned in two  
forms: with and without stop-codon (to allow fusion with C-terminal  
tag). The CDS has been directionally cloned using BD In-Fusion(TM)  
vector. Additional sequences in the clone: 'ACC' after SalI site  
and before 'ATG' to provide Kozak consensus sequence; 'GG' after  
last codon and before HindIII site to maintain reading frame.  
clone distribution: <http://bioinfo.clontech.com/orfclones>.  
FEATURES  
source 1..963 Location/Qualifiers  
/organism="Homo sapiens"

3

CDS

```

/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="GH00457X1.0"
/lab_host="BD Creator(TM) CDS library derived from MGC
collection"
/lab_host="DH5alpha T1 resistant"
/notes="Vector: pDNR-Dual"
1..963
/codon_start=1
/product="eukaryotic translation initiation factor 3,
subunit 4 delta, 44kDa"
/protein_id="AAP35535.1"
/db_xref="GI:30582617"
/translation="MPTGDFDSKPSWADQVEEGEDDKCVTSLLKGIPLANGTSP
PELLGAPLPPEPKVINGNIKTVEIKIDEDKKKIVTFRFETKSAVARRW
KRGNSERPDPGNVATTVSDVSWTFITSKEDLNCOEEDPMNLKQKIVSCIC
KEDHWTRCPYKDTLGPQKELAEQGLSTGEKELPGELEPVQATONKTGKVP
RGSARSGSMQPNRRADNATIRVNLSIEDRETDLELFRPFGSISRILAKDKT
GOSKGFAPISFHRREDARAIAGVSGFGDHLILNVMAKPSNL"
BASE COUNT      237 a      291 c      288 g      147 t
ORIGIN
Query Match      88.9%; Score 16; DB 9; Length 963;
Best Local Similarity 100.0%; Pred. No. 15;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 GGCTGGTGCACCTGT 16
      |||||
Db      127 GGCTGGTGCACCTGT 112

RESULT 19
BT007572/c      963 bp      mRNA      linear      SYN 13-MAY-2003
LOCUS      Synthetic construct Homo sapiens eukaryotic translation initiation
DEFINITION      factor 3, subunit 4 delta, 44kDa mRNA, partial cds.
ACCESSION      BT007572
VERSION      BT007572.1 GI:30583982
KEYWORDS      FLI CDNA.
SOURCE      Synthetic construct
ORGANISM      synthetic construct
REFERENCE      1 (bases 1 to 963)
AUTHORS      Kalnina,N., Chen,X., Rolfs,A., Halleck,A., Hines,L., Eisenstein,S.,
Koundinya,M., Raphael,J., Moreira,D., Kelley,T., Labaer,J., Lin,Y.,
Phelan,M., and Farmer,A.
JOURNAL      Cloning of human full-length CDSs in BD Creator(TM) System Donor
TITLE      Unpublished
vector
2 (bases 1 to 963)
Kalinina,N., Chen,X., Rolfs,A., Halleck,A., Hines,L., Eisenstein,S.,
Koundinya,M., Raphael,J., Moreira,D., Kelley,T., Labaer,J., Lin,Y.,
Phelan,M., and Farmer,A.
DIRECT SUBMISSION
Submitted (13-MAY-2003) BD Biosciences Clontech, 1020 East Meadow
Circle, Palo Alto, CA 94303, USA
COMMENT      This CDS clone is a part of a collection of human full length
expression clones generated by BD Biosciences Clontech and the
Harvard Institute of Proteomics. Each CDS has been cloned in two
forms: with and without stop-codon (to allow fusion with C-terminal
tag). The CDS has been directionally cloned using BD in-Fusion(TM)
cloning system between the SalI and HindIII sites of the pDNR-DUAL
vector. Additional sequences in the clone: 'ACC' after SalI site
and before 'ATG' to provide Kozak consensus sequence; 'GG' after
last codon and before HindIII site to maintain reading frame.
Clone distribution: http://bioinfo.clontech.com/orfclones.
location/Qualifiers
1..963
/organism="synthetic construct"
/mol_type="mRNA"
/db_xref="taxon:32630"
/clone="GH00457L1.0"
/clone_1ib="BD Creator(TM) CDS library derived from MGC

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CDS

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collection"
/lab_host="DH5alpha T1 resistant"
/notes="Vector: pDNR-Dual"
1..>963
/notes="Mutations: 962:Stop->Leu"
/codon_start=1
/transl_table=11
/product="Homo sapiens eukaryotic translation initiation
factor 3, subunit 4 delta, 44kDa"
/protein_id="AAP36240.1"
/db_xref="GI:30583983"
/translation="MPTGDFDSKPSWADQVEEGEDDKCVTSLLKGIPLANGTSP
PELLGAPLPPEPKVINGNIKTVEIKIDEDKKKIVTFRFETKSAVARRW
KRGNSERPDPGNVATTVSDVSWTFITSKEDLNCOEEDPMNLKQKIVSCIC
KEDHWTRCPYKDTLGPQKELAEQGLSTGEKELPGELEPVQATONKTGKVP
RGSARSGSMQPNRRADNATIRVNLSIEDRETDLELFRPFGSISRILAKDKT
GOSKGFAPISFHRREDARAIAGVSGFGDHLILNVMAKPSNL"
BASE COUNT      236 a      291 c      288 g      148 t
ORIGIN
Query Match      88.9%; Score 16; DB 12; Length 963;
Best Local Similarity 100.0%; Pred. No. 15;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 GGCTGGTGCACCTGT 16
      |||||
Db      127 GGCTGGTGCACCTGT 112

RESULT 20
104174/c      1041 bp      ss-DNA      linear      PAT 21-MAY-1993
LOCUS      Sequence 3 from Patent US 4707358.
DEFINITION
ACCESSION      104174
VERSION      104174.1 GI:268757
KEYWORDS
SOURCE      Unknown.
ORGANISM      Unknown.
REFERENCE      1 (bases 1 to 1041)
AUTHORS      Kieff,E., Tanner,J., Hummel,M. and Beisel,C.
JOURNAL      Vaccine against Epstein-Barr Virus
Patent: US 4707358-A 3 17-NOV-1987;
The University of Chicago; Chicago, IL
location/Qualifiers
1..1041
/organism="unknown"
BASE COUNT      295 a      383 c      199 g      164 t
ORIGIN
Query Match      88.9%; Score 16; DB 6; Length 1041;
Best Local Similarity 100.0%; Pred. No. 15;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 GGCTGGTGCACCTGT 16
      |||||
Db      444 GGCTGGTGCACCTGT 429

RESULT 21
AF020833/c      1103 bp      mRNA      linear      PRI 02-MAR-1999
LOCUS      Homo sapiens eukaryotic translation initiation factor 3 subunit
DEFINITION      (p42) mRNA, complete cds.
ACCESSION      AF020833
VERSION      AF020833.1 GI:2460199
KEYWORDS
SOURCE      Homo sapiens (human)
ORGANISM      Homo sapiens
REFERENCE      Bandyopadhyay,A. and Maitra,U.
1 (bases 1 to 1103)
Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo.
Bandyopadhyay,A. and Maitra,U.

```



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/codon_start=1
/product="eukaryotic translation initiation factor 3,
subunit 4 (delta, 44kD)"
/subunit_id="AAH00733.1"
/protein_id="AAH00733.1"
/db_xref="GI:12653883"
/translation="MPTGDFSKPSWADQVEEGEDDKCVTSLLKGIPLATGDSPE
PELLGAPLPPEKVEVINGIKVTEYKIDEDGKKFIVTFRIETRKASKAVARRNM
KKEGSEFPPGPNVATTTSDVSMFTFITSKEDLNCOEEDPMNKLKQKIVSCRIC
KDDHMTTRCPYDITGPMOKELAEOLGISTGKEXLPGLEPVQATONKTKGVPPSL
RGCASRGESMGPNNRRADNATIRTNLSDETRDLOELFRPFGSISIIYLAOKXTT
GOSKGFATISFRRREDAPALAGVSGFGDHLILNVEWAKPSTN"

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BASE COUNT      283 a      336 c      335 g      174 t
ORIGIN
Query Match      88.9%; Score 16; DB 9; Length 1128;
Best Local Similarity 100.0%; Pred. No. 15;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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OY      1 GGCTGGTGCACCTGT 16
      |||||
Db      178 GGCTGGTGCACCTGT 163

RESULT 24
BC008469/c      1138 bp      mRNA      linear      PRI 12-JUL-2001
LOCUS      BC008469
DEFINITION      Homo sapiens, eukaryotic translation initiation factor 3, subunit 4
(delta, 44kD), clone MGC:14741 IMAGE:4279770, mRNA, complete cds.
ACCESSION      BC008469
VERSION      BC008469.1 GI:14250113
KEYWORDS      MGC.
SOURCE      Homo sapiens (human)
ORGANISM      Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 1138)
Strausberg, R.
Direct Submission
Submitted (25-MAY-2001) National Institutes of Health, Mammalian
Gene Collection (MGC), Cancer Genomics Office, National Cancer
Institute, 31 Center Drive, Room 11A03, Bethesda, MD 20892-2590,
USA
NIH-MGC Project URL: http://mgc.nci.nih.gov
Contact: MGC help desk
Email: cgabbs-r@mail.nih.gov
Tissue Procurement: ATCC
CDNA Library Preparation: CLONTECH Laboratories, Inc.
DNA Library Arrayed by: The I.M.A.G.E. Consortium (LNL)
DNA Sequencing by: Sequencing Group at the Stanford Human Genome
Center, Stanford University School of Medicine, Stanford, CA 94305
Web site: http://www-shgc.stanford.edu
Contact: (Dickson, Mark) mcd@paxil.stanford.edu
Dickson, M., Schmutz, J., Grimwood, J., Rodriguez, A., and Myers,
R. M.

```

Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LNL at: <http://image.lnl.gov>  
Series: IRIL Plate: 21 Row: e Column: 24  
This clone was selected for full length sequencing because it passed the following selection criteria: Similarity but not identity to protein.

FEATURES  
source  
Location/Qualifiers.

```

1..1138
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="MGC:14741 IMAGE:4279770"
/issue_type="Brain, primitive neuroectodermal"
/clone_id="N1H_MGC_56"
/lab_host="DH10B"
/notes="Vector: pDNR-L1B"
44..1006

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CDS

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/codon_start=1
/product="eukaryotic translation initiation factor 3,
subunit 4 (delta, 44kD)"
/subunit_id="AAH08469.1"
/protein_id="AAH08469.1"
/db_xref="GI:14250114"
/translation="MPTGDFSKPSWADQVEEGEDDKCVTSLLKGIPLATGDSPE
PELLGAPLPPEKVEVINGIKVTEYKIDEDGKKFIVTFRIETRKASKAVARRNM
KKEGSEFPPGPNVATTTSDVSMFTFITSKEDLNCOEEDPMNKLKQKIVSCRIC
KDDHMTTRCPYDITGPMOKELAEOLGISTGKEXLPGLEPVQATONKTKGVPPSL
RGCASRGESMGPNNRRADNATIRTNLSDETRDLOELFRPFGSISIIYLAOKXTT
GOSKGFATISFRRREDAPALAGVSGFGDHLILNVEWAKPSTN"

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BASE COUNT      297 a      334 c      331 g      176 t
ORIGIN
Query Match      88.9%; Score 16; DB 9; Length 1138;
Best Local Similarity 100.0%; Pred. No. 15;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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```

OY      1 GGCTGGTGCACCTGT 16
      |||||
Db      170 GGCTGGTGCACCTGT 155

RESULT 25
BD063236/c      1142 bp      DNA      linear      PAT 27-AUG-2002
LOCUS      BD063236
DEFINITION      Secreted human proteins.
ACCESSION      BD063236
VERSION      BD063236.1 GI:22608839
KEYWORDS      JP 2001505783-A/11.
SOURCE      unidentified
ORGANISM      unidentified
unclassified.
1 (bases 1 to 1142)
Escobedo, J., Hu, Q., Garcia, P., Williams, L.T. and Kothakota, S.
Secreted human proteins
Patent: JP 2001505783-A 11 08-MAY-2001;
CHIRON CORP
PN JP 2001505783-A/11
PD 08-MAY-2001
PF 11-DEC-1997 JP 1998526977
PR 11-DEC-1996 US 60/032757
PI JAIME ESCOBEDO, QUINJUN HU, PABLO GARCIA, LEWIS T WILLIAMS PI
SRINIVAS KOTHA KOTA
PC C12N15/12, C12N15/62, C12N15/85, C12N5/10, C12N1/21, C07K14/47, PC
C07K16/18
CC Strandedness: Single;
CC Topology: Linear;
FH Key Location/Qualifiers.
1..1142
/organism="unidentified"
/mol_type="genomic DNA"
/db_xref="taxon:32644"

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FEATURES

source

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BASE COUNT      304 a      335 c      331 g      172 t
ORIGIN

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Query Match      88.9%; Score 16; DB 6; Length 1142;
Best Local Similarity 100.0%; Pred. No. 15;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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OY      1 GGCTGGTGCACCTGT 16
      |||||
Db      159 GGCTGGTGCACCTGT 144

RESULT 26
AF094850/c      1174 bp      mRNA      linear      PRI 23-MAR-2001
LOCUS      AF094850
DEFINITION      Homo sapiens eukaryotic translation initiation factor 3 subunit
p42/p44 mRNA, complete cds.
ACCESSION      AF094850
VERSION      AF094850.1 GI:10280561

```



KEYWORDS  
SOURCE  
ORGANISM  
REFERENCE  
AUTHORS  
TITLE  
JOURNAL  
REFERENCE  
AUTHORS  
TITLE  
JOURNAL  
FEATURES  
source

Homo sapiens (human)  
Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.  
1 (bases 1 to 1174)  
Chen, W., Blough, R. I., and Winkelman, J. C.  
Molecular cloning, genomic structure and chromosomal localization  
of a novel human RNA binding protein gene homologous to a tumor  
necrosis factor alpha inducible transcript in mouse  
Unpublished  
2 (bases 1 to 1174)  
Chen, W., Chu, Z., Blough, R. I., Liu, L., Hoppes, B. and  
Winkelman, J. C.  
Direct Submission  
Submitted (24-SEP-1998) Internal Medicine/Hematology-Oncology,  
University of Cincinnati College of Medicine, 231 Bethesda Ave.,  
Cincinnati, OH 45267-0508, USA  
Location/Qualifiers  
1. 1174  
/organism="Homo sapiens"  
/mol\_type="mRNA"  
/db\_xref="taxon:9606"  
/chromosome="19"  
/map="19p13.2"  
42. 1004  
/note="eIF3 p42/p44"  
/product="eukaryotic translation initiation factor 3  
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/db\_xref="GI:10280562"  
/translation="MPTGDFSKPSMADQVEBERGDDKCVTSLLKGIPIATDTSPE  
PELLPGAPLPPEPKVINGNTKTEYKIDDDGKKPKIVRFRLETRKASAVARRKM  
KRRNSBEPDPPGNNVAFTYSDVSMFTLSKEDLNQSEEDMNTLKKQKXISCRIC  
KEDWTRCPCYKDTLGMQKELABQVSLSTGEKPKFGELEFPQATNRTKGVYPSL  
RDGASRRGSMQPNRRADNATRVNLSDTEETDQLFRFGSISRIYLAKDKYT  
GSGKPAFISFHRERDAARAIAAGVSGFGYDHLINEMAKPSTN"

BASE COUNT 334 a 336 c 331 g 173 t

ORIGIN

Query Match 88.9%; Score 16; DB 9; Length 1174;  
Best Local Similarity 100.0%; Pred. No. 15;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGCTGGTGCACCTGT 16  
168 GGCTGGTGCACCTGT 153

Db

RESULT 27  
AF348496/c 2199 bp DNA linear PLN 21-AUG-2001  
LOCUS  
DEFINITION  
VERSION  
ACCESSION  
KEYWORDS  
SOURCE  
ORGANISM

Desmodemus pirkollei  
Desmodemus pirkollei  
Eukaryota; Viridiplantae; Chlorophyta; Chlorophyceae;  
Chlorococcales; Scenedesmeceae; Desmodemus.  
1 (bases 1 to 2199)  
Hegewald, E., Coesel, P. and Hegewald, P.  
A phytoplankton collection from Bali, with description of a new  
Desmodemus species (Chlorophyta, Scenedesmeceae)  
Unpublished  
2 (bases 1 to 2199)  
Hegewald, E., Coesel, P. and Hegewald, P.  
Direct Submission  
Submitted (13-FEB-2001) Institute of Chemistry and Dynamics of the  
Geosphere 6, Research Center Juelich, Juelich D-52425, Germany  
Location/Qualifiers  
1. 2199

REFERENCE  
AUTHORS  
TITLE  
JOURNAL  
FEATURES  
source

RNA  
BASE COUNT  
ORIGIN  
Query Match 88.9%; Score 16; DB 8; Length 2199;  
Best Local Similarity 100.0%; Pred. No. 14;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGCTGGTGCACCTGT 16  
1393 GGCTGGTGCACCTGT 1378

Db

RESULT 28  
AF172333/c 2556 bp mRNA linear VRL 29-AUG-1999  
LOCUS  
DEFINITION  
VERSION  
ACCESSION  
KEYWORDS  
SOURCE  
ORGANISM

Human herpesvirus 4 cell-line SNU-1103 major outer envelope  
glycoprotein gp350 mRNA, complete cds.  
AF172333.1 GI:5802484  
Human herpesvirus 4 (Epstein-Barr virus)  
Human herpesvirus 4  
Viruses; dsDNA viruses, no RNA stage; Herpesviridae;  
Gammaherpesvirinae; Lymphocryptovirus.  
1 (bases 1 to 2556)  
Lee, W.K., Kim, S.W., Sim, Y.S., Cho, S.G., Park, S.H., Kim, C.W. and  
Park, J.G.  
B-lymphoblastoid cell lines from cancer patients  
In Vitro Cell. Dev. Biol. Anim. 34 (2), 97-100 (1998)  
9542645  
2 (bases 1 to 2556)  
Chang, S.H., Kim, S.H., Lee, W.K., Kim, H.J., Choi, S.H., Park, J.H.,  
Jang, H.S., Chung, G.H., Kwon, T.H., Kim, D.H., Yang, M.S. and Jang, Y.S.  
Cloning and analysis of the Epstein-Barr virus glycoprotein 350  
genes  
Mol. Cells 8 (5), 585-593 (1998)  
99072166  
MEDLINE  
PUBMED  
REFERENCE  
AUTHORS  
TITLE  
JOURNAL  
FEATURES  
source

3 (bases 1 to 2556)  
Chang, S.H., Kim, S.H., Lee, W.K., Kim, H.J., Choi, S.H., Park, J.H.,  
Jang, H.S., Chung, G.H., Kwon, T.H., Kim, D.H., Yang, M.S. and Jang, Y.S.  
Direct Submission  
Submitted (27-JUL-1999) Faculty of Biological Sciences, Chonbuk  
National University, 664-14, Dukjin-dong 1-Ka, Dukjin-Ku, Chonju  
561-756, Korea  
Location/Qualifiers  
1. 2556  
/organism="Human herpesvirus 4"  
/mol\_type="mRNA"  
/db\_xref="taxon:10376"  
/cell\_line="SNU-1103"  
1. 2556  
/codon\_start=1  
/product="major outer envelope glycoprotein gp350"  
/protein\_id="AA051698.1"  
/translation="MEALLVCOYTIQSLIQLTRDDPGFVMEILEFPYPACNVCTA  
DNATINPDGKQOLNDFQOLPPTAVYVOPRGAFGSENAITLFLLELGAEL  
ALTRSKKPIVVTGGEBOVLESDVYFOVFGTGMCHHMEMONPVLLIPTVYI  
KNDNCSTITAVRAQGDVTLPLSLPSADSNYSYKTMAGNIDIECTDEDEI  
SOVLPGDNFNITCSGIESHPVSGGLITSIPVATIPGIGTAYISRLTPRPSRSLG  
NNSILVYFSGNGPKASGDYCIQSNIVSDEIPASQDMPTTDTITVGDNAVYSP  
NVTSEDAANPVTATVAFAMPNNTETDFPKWTLSGTPSGCENISGAFASNTPTIT  
VSLGAPMTLITRTATNATTTTHKIVRSKAPESYTSPTLNTTGFAPNPTTGAPS  
GTHVPNTLTAPASTGPTVSTADVTSPPTAGTSGASPVNPSGPRNGTESKADMTS  
PTSAVTTTPPNAITSPPTAVTTTPPNAITSLTGKTSPTSAVTTTPPNAITSPYGTSPQ  
ANTNHTIGTSTSPVTSPPKNAITSVTTGHNITSSSTSSMSLSETSISSETLSPST



SDNSTSHMPLTSAHPTGENTITVOYTPASTSTHNVSTSSPAPRPCTTQASGPNSSST  
TEKPEVAVTTCPTPKNATSPAPSGOKTAVPTVSTGKANSITGKTTGAGRTS  
TPTPDYGDSTTPRTYATVTLPTSTSKLRPMTFTSPPTTAQATVPVPSQOP  
RNSMLVQMASLAVTLTLLLMADACAFRNLSTSHYTTTPPYDAETVY"

BASE COUNT 713 a 798 c 532 g 513 t

ORIGIN

Query Match 88.9%; Score 16; DB 14; Length 2556;  
Best Local Similarity 100.0%; Pred. No. 14;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGCTGGTGCACCTGT 16  
|||||

Db 1956 GGCTGGTGCACCTGT 1941

RESULT 29  
AF172332 2661 bp mRNA linear VRL 29-AUG-1999  
LOCUS Human herpesvirus 4 cell-line SNU-20 major outer envelope  
DEFINITION glycoprotein gp350 mRNA, complete cds.  
ACCESSION AF172332  
VERSION AF172332.1 GI:5802482  
KEYWORDS Human herpesvirus 4 (Epstein-Barr virus)  
SOURCE Viruses; dsDNA viruses, no RNA stage; Herpesviridae;  
ORGANISM Gammaherpesvirinae; Lymphocryptovirus.  
REFERENCE 1 (bases 1 to 2661)  
AUTHORS Lee, W.K., Kim, S.M., Sim, Y.S., Cho, S.G., Park, S.H., Kim, C.W. and  
Park, J.G.  
TITLE B-lymphoblastoid cell lines from cancer patients  
JOURNAL In Vitro Cell. Dev. Biol. Anim. 34 (2), 97-100 (1998)  
MEDLINE 98203772  
PUBMED 9542645  
REFERENCE 2 (bases 1 to 2661)  
AUTHORS Chang, S.H., Kim, S.H., Lee, W.K., Kim, H.J., Choi, S.H., Park, J.H.,  
Jang, H.S., Chung, G.H., Kwon, T.H., Kim, D.H., Yang, M.S. and Jang, Y.S.  
TITLE Cloning and analysis of the Epstein-Barr virus glycoprotein 350  
JOURNAL Mol. Cells 8 (5), 585-593 (1998)  
MEDLINE 99072166  
PUBMED 9856346  
REFERENCE 3 (bases 1 to 2661)  
AUTHORS Chang, S.H., Kim, S.H., Lee, W.K., Kim, H.J., Choi, S.H., Park, J.H.,  
Jang, H.S., Chung, G.H., Kwon, T.H., Kim, D.H., Yang, M.S. and Jang, Y.S.  
TITLE Direct Submission  
JOURNAL Submitted (27-JUL-1999) Faculty of Biological Sciences, Chonbuk  
National University, 664-14, Dujin-dong 1-Ka, Dujin-Ku, Chonju  
561 - 756, Korea  
FEATURES  
source Location/Qualifiers  
1..2661  
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/db\_xref="taxon:10376"  
/cell\_line="SNU-20"  
1..2661  
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DVNATINPDVGGKKKLNLDLGLTPHTKAVYOPRGAFGSEKATNLFLELLAGSEL  
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KNDNCSTNITAVRAQGLDVTLPISLPSADSNFSVKTMLGNIDIECMEDEI  
SOVLPGDNFNITCSGYESHVPSGGLITSLPSVATPIPGGYASRLRTPVRSRLG  
NNSILVYFSGNGPKASGGDYCIOSNIVSDEI PASQDMPNTTDTTYGDNAYSVP  
NWSIEDANPNVTVTAFAMPNNTETDFCKWTLNSGTPSGCENTISGAPASNTEDIT  
VSGLGAPKTLITRTATNATTTTHKVIKASBESTTPTLNTTTPAAPTNTTGLPS  
STHVPNTLAPASTGPTVSTADVTSPTPAGTSGASPVTPRSPRNGESKADMTS  
PTSAVTPTPNATSPPTAVTPTPNATSPGLGTSPTSAVTPTPNATSPPTAVTPT  
PNATIPTLGTSPTSAVTPTPNATSPVAGETSPKANTNHTLGGTSSTPVVTSPPKN  
ATSAVTGQHNITSSSTSSMSLRPSSISFTLSPSTSDNSMSHMLTSAHPGCEMIT

QVTPASTSTHNVSTSSPAPRPCTTQASGPNSSSTKRGVAVTTCPTPKNATSPQA  
PSGOKTAVPTVSTGKANSITGKTTGAGRTSPTPDYGDSTTPRTYATVTL  
LPTSTSKLRPMTFTSPPTTAQATVPVPSQOPRFSNLSMLVQMASLAVTLTLL  
LVMADACAFRNLSTSHYTTTPPYDAETVY"

BASE COUNT 752 a 844 c 542 g 523 t

ORIGIN

Query Match 88.9%; Score 16; DB 14; Length 2661;  
Best Local Similarity 100.0%; Pred. No. 14;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGCTGGTGCACCTGT 16  
|||||

Db 2061 GGCTGGTGCACCTGT 2046

RESULT 30  
EBVBLF1A/C 2661 bp DNA linear VRL 23-OCT-1992  
LOCUS Epstein-Barr virus BLF1 gene for glycoprotein 350/220.  
DEFINITION X67776  
ACCESSION X67776  
VERSION X67776.1 GI:59163  
KEYWORDS glycoprotein 350/220.  
SOURCE Human herpesvirus 4 (Epstein-Barr virus)  
ORGANISM Viruses; dsDNA viruses, no RNA stage; Herpesviridae;  
REFERENCE 1 (bases 1 to 2661)  
AUTHORS Klein, K. and Mueller-Lantzsch, N.  
TITLE Sequences of the membrane proteins gp 350/220 and p140 of  
Epstein-Barr virus type-B (P3HR1)  
JOURNAL Nucleic Acids Res. 2 (bases 1 to 2661)  
AUTHORS Klein, K.  
TITLE Direct Submission  
JOURNAL Submitted (21-OCT-1992) K. Klein, Inst f Med Mikrobiologie u  
Hygiene, Abteilung Virologie, Universitaetsklinikum des Saarlandes,  
6650 Homburg/Saar, FRG  
FEATURES  
source Location/Qualifiers  
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/protein\_id="CAA47986.1"  
/db\_xref="GI:59164"  
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DVNATINPDVGGKKKLNLDLGLTPHTKAVYOPRGAFGSEKATNLFLELLAGSEL  
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KNDNCSTNITAVRAQGLDVTLPISLPSADSNFSVKTMLGNIDIECMEDEI  
SOVLPGDNFNITCSGYESHVPSGGLITSLPSVATPIPGGYASRLRTPVRSRLG  
NNSILVYFSGNGPKASGGDYCIOSNIVSDEI PASQDMPNTTDTTYGDNAYSVP  
NWSIEDANPNVTVTAFAMPNNTETDFCKWTLNSGTPSGCENTISGAPASNTEDIT  
VSGLGAPKTLITRTATNATTTTHKVIKASBESTTPTLNTTTPAAPTNTTGLPS  
STHVPNTLAPASTGPTVSTADVTSPTPAGTSGASPVTPRSPRNGESKADMTS  
PTSAVTPTPNATSPPTAVTPTPNATSPGLGTSPTSAVTPTPNATSPPTAVTPT  
PNATIPTLGTSPTSAVTPTPNATSPVAGETSPKANTNHTLGGTSSTPVVTSPPKN  
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Best Local Similarity 100.0%; Pred. No. 14;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGCTGGTGCACCTGT 16  
Db 2061 GGCTGGTGCACCTGT 2046

RESULT 31  
EBVBLF1/c 2663 bp DNA linear VRL 09-JAN-1998  
LOCUS Epstein-Barr virus BLF1 gene..  
X99106  
VERSION X99106.1 GI:2769559  
KEYWORDS BLF1 gene; gp340.  
SOURCE Human herpesvirus 4 (Epstein-Barr virus)  
ORGANISM Human herpesvirus 4  
Viruses; dsDNA viruses, no RNA stage; Herpesviridae;  
Gammaherpesvirinae; Lymphocryptovirus.

REFERENCE  
AUTHORS Mackett, M., Cox, C., Pepper, S. D., Lees, J. F., Naylor, B. A.,  
Wedderburn, N. and Arrand, J. R.  
TITLE Immunisation of common marmosets with vaccinia virus expressing  
Epstein-Barr virus (EBV) gp340 and challenge with EBV  
J. Med. Virol. 50 (3), 263-271 (1996)  
MEDLINE 97082049  
PUBMED 8923292  
REFERENCE 2 (bases 1 to 2663)  
AUTHORS Pepper, S. D. V.  
TITLE Direct Submission  
JOURNAL Submitted (04-JUL-1996) S.D.V. Pepper, Paterson Institute for  
Cancer Research, Molecular Biology, Wilmslow Rd, Wilington,  
Manchester M20 9BX, UK

FEATURES  
source  
1..2663  
/organism="Human herpesvirus 4"  
/mol\_type="genomic DNA"  
/strain="M81"  
/db\_xref="taxon:10376"  
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KVNCSNTITAVRAQGLVTLPLSPSAQDSNPFVKOMLGNEDIECIDDEI  
SOYLPQDNKNTCSGESHPVSGGIIITISPVATPIRGYVYSLRLTVPYRFLG  
NNSILVYISGNGPKASGDYCIQSNIVFSDIIPASDMPNTTIDITYVDNAVYVP  
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VSLGAPKTLITRTATNATTTTKVIFSKAPESITTSPLNTGTGADPNITGLPS  
STHVPNLTPASTGPTVADYVTPASTGASVTPSPSPRODGTESKADMTS  
PTSAVTPTPNGTSPPTAMTTPPNATSPRTIGTSPSAATTPTPNATSPPAVTP  
PNATSPVGTSPQANATNTILGISTPVTVPPTPKATSDVTGGHNRSSSTSSNS  
LRPSSIPETTSHPPLTSAHPGTGENTQVTPASISTHNVSTSSPAAPRTTSQASGP  
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GARTSTPTDYGDDSTTPRYNATYLPSPSSSKLRPMWTFSPPTTAQATVFP  
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BASE COUNT 755 a 829 c 552 g 527 t  
ORIGIN

Query Match 88.9%; Score 16; DB 14; Length 2663;  
Best Local Similarity 100.0%; Pred. No. 14;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 1 GGCTGGTGCACCTGT 16

Db 2036 GGCTGGTGCACCTGT 2021

RESULT 32  
E00513/c 2721 bp DNA linear PAT 29-SEP-1997  
LOCUS Genomic DNA encoding Epstein-Barr virus gp 220/200.  
E00513  
VERSION E00513.1 GI:2168792  
KEYWORDS JP 1985232094-A/1.  
SOURCE Human herpesvirus 4 (Epstein-Barr virus)  
ORGANISM Human herpesvirus 4  
Viruses; dsDNA viruses, no RNA stage; Herpesviridae;  
Gammaherpesvirinae; Lymphocryptovirus.

REFERENCE  
AUTHORS Ericot, K., Jieroom, T., Mearti, F. and Kuristofua, B.  
TITLE VACCINE TO EPSTEIN BAR VIRUS  
JOURNAL Patent: JP 1985232094-A 1 18-NOV-1985;  
UNIV CHICAGO:THE

COMMENT  
OS Epstein-Barr virus  
PN JP 1985232094-A/1  
PD 18-NOV-1985  
PF 30-JAN-1985 JP 1985014606  
PR 30-JAN-1984 US 84 575352, 23-JUL-1984 US 84 633558 PI  
ERIOSTUTO KILFU, JIEROOMU TANNA, MEARTI FUNMERU, PI KURISTOFUUA  
BEISERU

PC C12N15/00, A61K39/245, C07H21/04, C07K13/00, C12N1/00//C12P21/02,  
PC C12N1/00,  
PC C12R1:19);  
CC strandedness: Double;  
CC topology: Linear;  
CC hypothetical: No;  
CC anti-sense: No;  
FH key  
FT Location/Qualifiers  
CDS 1..>2721  
/product="gp 220/200".  
Location/Qualifiers  
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/mol\_type="genomic DNA"  
/db\_xref="taxon:10376"

BASE COUNT 762 a 876 c 557 g 526 t  
ORIGIN

Query Match 88.9%; Score 16; DB 6; Length 2721;  
Best Local Similarity 100.0%; Pred. No. 14;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGCTGGTGCACCTGT 16  
Db 2124 GGCTGGTGCACCTGT 2109

RESULT 33  
HS4GPR340A 3210 bp DNA linear VRL 15-FEB-2001  
LOCUS Epstein-Barr virus glycoprotein 340 (BLF1) and BLF2 genes,  
complete cds's, and BLRF3 gene, first exon.  
H54GPR340A  
VERSION 107922.1 GI:291519  
KEYWORDS antigen; glycoprotein 340; gp340; viral antigen.  
SOURCE Human herpesvirus 4 (Epstein-Barr virus)  
ORGANISM Human herpesvirus 4  
Viruses; dsDNA viruses, no RNA stage; Herpesviridae;  
Gammaherpesvirinae; Lymphocryptovirus.

REFERENCE  
AUTHORS Lees, J. F., Arrand, J. B., Pepper, S. D., Stewart, J. P., Mackett, M. and  
Arrand, J. R.  
TITLE The Epstein-Barr virus candidate vaccine antigen gp340/220 is  
highly conserved between virus types A and B  
JOURNAL Virology 195 (2), 578-586 (1993)

MEDLINE 93331716  
PUBMED 8393237  
FEATURES  
source

Location/Qualifiers  
1..3210  
/organism="Human herpesvirus 4"  
/mol\_type="genomic DNA"  
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/db\_xref="taxon:10376"  
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complement(1..2661)  
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/db\_xref="GI:291520"  
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NSVILYFSGNGPRASGSDYCIQSNIVSDEIPASQDMFTNTDITTYGDNATYVP  
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SEERADAHLDSSQKRRKRRVDDAGSNAPQHVPPQDILHGRALYRFLDLRR  
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BASE COUNT 659 a 678 c 1001 g 872 t

ORIGIN

Query Match 88.9%; Score 16; DB 14; Length 3210;  
Best Local Similarity 100.0%; Pred. No. 14;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGCTGGTGCACCTGT 16  
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Db 601 GGCTGGTGCACCTGT 616

RESULT 34  
LOCUS A11128 3400 bp DNA linear PAT 07-DEC-1993  
DEFINITION Synthetic nucleotide sequence of the leftward reading frame of the  
Bam I-fragment encoding gp 250/350.  
ACCESSION A11128

VERSION A11128.1 GI:490970  
KEYWORDS  
SOURCE  
ORGANISM  
REFERENCE  
AUTHORS  
TITLE

JOURNAL  
Wolfe, H. J.  
DNA sequences of the EBV genome, recombinant DNA molecules,  
processes for producing EBV-related antigens, diagnostic  
compositions and pharmaceutical compositions containing said  
antigens  
Patent: EP 0173254-A 3 05-MAR-1986;  
Wolfe, Hans Joachim

FEATURES  
source

Location/Qualifiers  
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/organism="synthetic construct"  
/mol\_type="genomic DNA"  
/db\_xref="taxon:32630"  
556..3279  
/note="Leftward reading frame of the Bam I-fragment  
encoding gp 250/350."  
/codon\_start=1  
/transl\_table=1  
/protein\_id="CA00938.1"  
/db\_xref="GI:490971"  
/translation="MEALLVCQYITQSLIHLTGDDPGFNFVEIEFPYPACNVCTA  
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BASE COUNT 904 a 1060 c 732 g 704 t

ORIGIN

Query Match 88.9%; Score 16; DB 6; Length 3400;  
Best Local Similarity 100.0%; Pred. No. 14;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGCTGGTGCACCTGT 16  
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Db 2679 GGCTGGTGCACCTGT 2664

RESULT 35  
LOCUS E01006 3400 bp DNA linear PAT 29-SEP-1997  
DEFINITION CDNA encoding p350 of Epstein-Barr virus.  
ACCESSION E01006  
VERSION E01006.1 GI:2169265  
KEYWORDS JP 1986257188-A/2.  
SOURCE Human herpesvirus 4 (Epstein-Barr virus)  
ORGANISM Human herpesvirus 4  
Viruses; deDNA viruses, no RNA stage; Herpesviridae;  
Gammaherpesvirinae; Lymphocryptovirus.  
Hansu, Y. B.  
1 (bases 1 to 3400)

REFERENCE  
AUTHORS  
TITLE

JOURNAL  
HANSU YOTSUTO BUORUFU  
OS Epstein-Barr virus  
PN JP 1986257188-A/2

PD 14-NOV-1986  
 PF 23-AUG-1985 JP 1985185661  
 PR 23-AUG-1984 EP 84 84110089, 23-AUG-1984 EP 84 84110090 PI  
 HANSU YOTSUTU BUORUFU  
 PC C12N15/00.A6IK39/245.C07H21/04.C07K13/00.C12N1/00.G0IN33/569,  
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 CC topology: Linear;  
 CC hypothetical: No;  
 CC anti-sense: No;  
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 FT CDS 556..3279  
 FT 3280..3400 /product='p350 Epstein-Barr virus' FT 3'UTR  
 FT polyA\_site 3284..3291.  
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 BASE COUNT 904 a 1060 c 732 g 704 t  
 ORIGIN

Query Match 88.9%; Score 16; DB 6; Length 3400;  
 Best Local Similarity 100.0%; Pred. No. 14;  
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 GGCTGTGTCACTGT 16  
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 Db 2679 GGCTGTGTCACTGT 2664

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 Job time : 503.975 secs

GenCore version 5.1.6  
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OM nucleic - nucleic search, using sw model

Run on: August 14, 2003, 21:32:41 ; Search time 547.75 Seconds  
(without alignments)  
1493.734 Million cell updates/sec

Title: US-10-074-620-2  
Perfect score: 20  
Sequence: 1 ccttaggaggaacagtcgcc 20

Scoring table: OLIGO\_NUC  
Gapop 60.0 , Gapext 60.0

Searched: 2888711 seqs, 2045481386 residues

Word size : 0

Total number of hits satisfying chosen parameters: 5777422

Minimum DB seq length: 0  
Maximum DB seq length: 2000000000

Post-processing: Listing first 120 summaries

Database :

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1: gb\_ba:\*  
2: gb\_hcg:\*  
3: gb\_in:\*  
4: gb\_om:\*  
5: gb\_ov:\*  
6: gb\_pac:\*  
7: gb\_ph:\*  
8: gb\_pl:\*  
9: gb\_pr:\*  
10: gb\_ro:\*  
11: gb\_sy:\*  
12: gb\_sy:\*  
13: gb\_un:\*  
14: gb\_vl:\*  
15: em\_ba:\*  
16: em\_fun:\*  
17: em\_hum:\*  
18: em\_in:\*  
19: em\_mu:\*  
20: em\_or:\*  
21: em\_or:\*  
22: em\_ov:\*  
23: em\_pac:\*  
24: em\_ph:\*  
25: em\_pl:\*  
26: em\_ro:\*  
27: em\_scs:\*  
28: em\_un:\*  
29: em\_vl:\*  
30: em\_hcg\_hum:\*  
31: em\_hcg\_inv:\*  
32: em\_hcg\_other:\*  
33: em\_hcg\_mus:\*  
34: em\_hcg\_pin:\*  
35: em\_hcg\_rod:\*  
36: em\_hcg\_mam:\*  
37: em\_hcg\_vrt:\*  
38: em\_sy:\*  
39: em\_hgo\_hum:\*  
40: em\_hgo\_mus:\*  
41: em\_hgo\_other:\*

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,  
and is derived by analysis of the total score distribution.

# SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	20	100.0	20	6	AX52237 Sequence
2	20	100.0	20	6	I26103 Sequence 29
3	20	100.0	1041	6	I04174 Sequence 3
4	20	100.0	2663	14	EBVBLFL1 X99106 Epstein-Bar
5	20	100.0	2721	6	E00513 Genomic DNA
6	20	100.0	3400	6	E11128 Synthetic n
7	20	100.0	3400	6	E01006 cDNA encodi
8	20	100.0	3833	6	AR049357 Sequence
9	20	100.0	5019	6	A11178 Synthetic n
10	20	100.0	5019	6	E01007 DNA sequenc
11	20	100.0	5931	6	AR233080 Sequence
12	20	100.0	5931	14	HS4ENVGP M10593 Epstein Bar
13	20	100.0	171823	14	HHV507799 Human her
14	20	100.0	172281	14	EBV V01555 Epstein-Bar
15	20	100.0	184113	14	HS4B958RAJ M80517 Epstein-Bar
16	17	85.0	163473	2	AC142486 Rattus no
17	17	85.0	193466	2	AC113548 AC142486
18	16	80.0	204	11	G20457 Mus muscu
19	16	80.0	1555	8	AY079380 Arabidops
20	16	80.0	1767	8	AY039920 Arabidops
21	16	80.0	2376	6	AX151433 Sequence
22	16	80.0	6203	6	AB002380 Human mRN
23	16	80.0	9501	9	AF180681 Homo sapi
24	16	80.0	16529	9	AF000271 Homo sapi
25	16	80.0	32813	9	AF520762 Homo sapi
26	16	80.0	41308	9	AC005203 Homo sapi
27	16	80.0	68589	2	AC087682 Homo sapi
28	16	80.0	76850	9	AC003109 Human DNA
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34	16	80.0	100000	9	AP000180 Homo sapi
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36	16	80.0	109741	8	AC004005 Arabidops
37	16	80.0	115293	9	AC127460 Homo sapi
38	16	80.0	117134	9	AL157904 Human DNA
39	16	80.0	120836	9	AC097457 Homo sapi
40	16	80.0	126626	9	AC104843 Homo sapi
41	16	80.0	137557	9	AC005909 Homo sapi
42	16	80.0	144234	2	AP002425 Homo sapi
43	16	80.0	144279	2	AP001570 Homo sapi
44	16	80.0	151741	2	AC018609 Mus muscu
45	16	80.0	153026	9	AL1391280 Human DNA
46	16	80.0	158621	2	AC092951 Homo sapi
47	16	80.0	160133	2	AC025140 Homo sapi
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63	16	80.0	192699	2	AC111102 Mus muscu
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65	16	80.0	196545	9	AC113390 Homo sapi

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67	16	80.0	206246	2	AL606485	AL606485 Homo sapi
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69	16	80.0	216972	9	AC067819	AC067819 Homo sapi
70	16	80.0	217586	2	AC016076	AC016076 Homo sapi
71	16	80.0	229910	2	AC135698	AC135698 Rattus no
72	16	80.0	229296	2	BX323549	BX323549 Mus muscu
73	16	80.0	257734	2	BX510318	BX510318 Mus muscu
74	16	80.0	261391	2	AC094647	AC094647 Rattus no
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81	15	75.0	305	1	AB106218	AB106218 Alteromon
82	15	75.0	654	11	BV013986	BV013986 S212P6774
83	15	75.0	702	11	BV030337	BV030337 S212P6206
84	15	75.0	713	9	HUMY287H02	HUMY287H02 Homo sapi
85	15	75.0	728	11	BV056104	BV056104 S212P6629
86	15	75.0	762	6	AX463524	AX463524 Sequence
87	15	75.0	1267	3	AK117030	AK117030 Ciona int
88	15	75.0	1546	3	AB058682	AB058682 Ciona int
89	15	75.0	1619	5	AB093327	AB093327 Silurana
90	15	75.0	1648	5	AB093328	AB093328 Silurana
91	15	75.0	1691	6	AX544370	AX544370 Sequence
92	15	75.0	2171	5	DYGOB65A	DYGOB65A Discomyge o
93	15	75.0	2556	14	AF172333	AF172333 Human her
94	15	75.0	2661	14	AF172332	AF172332 Human her
95	15	75.0	2661	14	EBVHLR1A	X67776 Epstein-Bar
96	15	75.0	2702	1	EFPPD1GNS	X96978 E. faecalis
97	15	75.0	3178	5	AF034606	AF034606 Dario rer
98	15	75.0	4043	10	HS4GP340A	LO7922 Epstein-Bar
99	15	75.0	4988	14	HS4GP340B	AF345864 Mus muscu
100	15	75.0	5207	10	AC006119	LO7923 Epstein-Bar
101	15	75.0	8647	10	AL669922	AC006119 Mus muscu
102	15	75.0	8797	10	AF028784	AL669922 Mouse DNA
103	15	75.0	16313	1	AE000879	AF028784 Rattus no
104	15	75.0	33106	10	AL928961	AE000879 Methanoba
105	15	75.0	46681	9	AC094009	AL928961 Mouse DNA
106	15	75.0	53572	2	AC100004	AC094009 Homo sapi
107	15	75.0	56944	2	AC100526	AC100004 Mus muscu
108	15	75.0	59905	2	AC100985	AC100526 Mus muscu
109	15	75.0	64150	9	AL161778	AC100985 Mus muscu
110	15	75.0	64739	2	AC101271	AL161778 Human DNA
111	15	75.0	68209	9	AL357275	AC101271 Mus muscu
112	15	75.0	70855	2	AC101491	AL357275 Human DNA
113	15	75.0	71425	2	AC098994	AC101491 Mus muscu
114	15	75.0	72274	2	AC145201	AC098994 Mus muscu
115	15	75.0	77588	2	AC018015	AC145201 Takifugu
116	15	75.0	79899	2	AC024620	AC018015 Drosophila
117	15	75.0	81862	9	AL133318	AC024620 Homo sapi
118	15	75.0	82369	10	AL929144	AL133318 Homo sapi
119	15	75.0	88763	2	AC139550	AL929144 Mouse DNA
120	15	75.0				AC139550 Homo sapi

## ALIGNMENTS

RESULT 1	AX522237	20 bp	DNA	linear	PAT 24-OCT-2002
LOCUS	AX522237	Sequence 2 from Patent WO02064842.			
DEFINITION	AX522237				
ACCESSION	AX522237.1	GI:24411115			
VERSION					
KEYWORDS					
SOURCE	Human herpesvirus 4 (Epstein-Barr virus)				
ORGANISM	Human herpesvirus 4				
REFERENCE	Witte, D.P. and Groen, P.A.				
AUTHORS					

TITLE Quantitative epstein barr virus per rapid assay  
JOURNAL Patent: WO 02064842-A 2 22-AUG-2002;  
CHILDREN'S HOSPITAL RESEARCH FOUNDATION (US)  
FEATURES Location/Qualifiers  
SOURCE 1.20  
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BASE COUNT 6 a 6 c 5 g 3 t

Query Match 100.0%; Score 20; DB 6; Length 20;  
Best Local Similarity 100.0%; Pred. No. 3.8;  
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CCTTAGGAGAACAGTCCC 20  
Db 1 CCTTAGGAGAACAGTCCC 20

RESULT 2  
LOCUS 126103 20 bp DNA linear PAT 07-OCT-1996  
DEFINITION Sequence 29 from patent US 5556772.  
ACCESSION 126103  
VERSION 126103.1 GI:1605973  
KEYWORDS  
SOURCE Unknown.  
ORGANISM Unclassified.  
REFERENCE 1 (bases 1 to 20)  
AUTHORS Sorge, J.A. and Mullinax, R.L.  
TITLE Polymerase compositions and uses thereof  
JOURNAL Patent: US 5556772-A 29 17-SEP-1996;  
FEATURES Location/Qualifiers  
SOURCE 1.20  
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BASE COUNT 6 a 6 c 5 g 3 t  
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Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CCTTAGGAGAACAGTCCC 20  
Db 1 CCTTAGGAGAACAGTCCC 20

RESULT 3  
LOCUS 104174 1041 bp ss-DNA linear PAT 21-MAY-1993  
DEFINITION Sequence 3 from Patent US 4707358.  
ACCESSION 104174  
VERSION 104174.1 GI:268757  
KEYWORDS  
SOURCE Unknown.  
ORGANISM Unknown.  
REFERENCE Unclassified.  
1 (bases 1 to 1041)  
AUTHORS Kieff, E., Tanner, J., Hummel, M. and Beisel, C.  
TITLE Vaccine against Epstein-Barr Virus  
JOURNAL Patent: US 4707358-A 3 17-NOV-1987;  
The University of Chicago, Chicago, IL  
FEATURES Location/Qualifiers  
SOURCE 1.1041  
/organism="unknown"

BASE COUNT 295 a 383 c 199 g 164 t  
Query Match 100.0%; Score 20; DB 6; Length 1041;  
Best Local Similarity 100.0%; Pred. No. 1.6;  
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CCTTAGAGAGAACAGTCCC 20  
 DB 206 CCTTAGAGAGAACAGTCCC 225

RESULT 4  
 LOCUS EBVBLF1 2663 bp DNA linear VRL 09-JAN-1998  
 DEFINITION Epstein-Barr virus BLF1 gene.  
 ACCESSION X99106  
 VERSION X99106.1 GI:2769559  
 KEYWORDS BLF1 gene; gp340.  
 SOURCE Human herpesvirus 4 (Epstein-Barr virus)  
 ORGANISM Human herpesvirus 4  
 Viruses; dsDNA viruses, no RNA stage; Herpesviridae; Gammaherpesvirinae; Lymphocryptovirus.

REFERENCE  
 1 Mackeiz,M., Cox,C., Pepper,S.D., Lees,J.F., Naylor,B.A., Wedderburn,N. and Arrand,J.R.  
 Immunisation of common marmosets with vaccinia virus expressing Epstein-Barr virus (EBV) gp340 and challenge with EBV  
 J. Med. Virol. 50 (3), 263-271 (1996)  
 PUBMED 8923292

2 (bases 1 to 2663)  
 Pepper,S.D.V.  
 Direct Submission  
 Submitted (04-JUL-1996) S.D.V. Pepper, Paterson Institute for Cancer Research, Molecular Biology, Wilmslow Rd, Withington, Manchester M20 9BX, UK

FEATURES  
 source 1.2663  
 Location/Qualifiers  
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 /db\_xref="taxon:10376"  
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 KMDNCSNTITAVVRAQGLDVLPLSLPSAODSNFSPRTOMLGNRPYDI  
 SOVLPGDNKFNITGSGYSHVBSGGLITSTSPVATPIPTGTAYASRLTPRVSRLG  
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Query Match 100.0%; Score 20; DB 14; Length 2663;  
 Best Local Similarity 100.0%; Pred. No. 1.3;  
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CCTTAGAGAGAACAGTCCC 20  
 DB 1825 CCTTAGAGAGAACAGTCCC 1844

RESULT 5  
 LOCUS E00513 2721 bp DNA linear PAT 29-SEP-1997  
 DEFINITION Genomic DNA encoding Epstein-Barr virus gp 220/200.  
 ACCESSION E00513  
 VERSION E00513.1 GI:2168792  
 KEYWORDS JP 1985232094-A/1.  
 SOURCE Human herpesvirus 4 (Epstein-Barr virus)  
 ORGANISM Human herpesvirus 4  
 Viruses; dsDNA viruses, no RNA stage; Herpesviridae; Gammaherpesvirinae; Lymphocryptovirus.

REFERENCE  
 1 (bases 1 to 2721)  
 Briotauto, K., Jieroomu, T., Mearli, F. and Kurisutofuaa, B.  
 VACCINE TO EPSTEIN BAR VIRUS  
 Patent: JP 1985232094-A 1 18-NOV-1985;  
 UNIV CHICAGO:THE

COMMENT  
 OS Epstein-Barr virus  
 PN JP 1985232094-A/1  
 PD 18-NOV-1985  
 PE 30-JAN-1985 JP 1985014606  
 PR 30-JAN-1984 US 84 575352, 23-JUL-1984 US 84 633558 PI  
 ERIOTSUTO KIIFU, JIEROOMU TANNA, MEARLI FUNMERU, PI KURISOTOFUAA  
 BEISERU

PC C12N15/00,A61K39/245,C07H21/04,C07K13/00,C12N1/00//C12P21/02,  
 PC C12N1/00,  
 PC C12R1.19),  
 CC strandedness: Double;  
 CC topology: Linear;  
 CC hypothetical: No;  
 CC anti-sense: No;  
 FH Key Location/Qualifiers  
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 /product="gp 220/200".  
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 ORIGIN

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 Best Local Similarity 100.0%; Pred. No. 1.3;  
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CCTTAGAGAGAACAGTCCC 20  
 DB 1886 CCTTAGAGAGAACAGTCCC 1905

RESULT 6  
 LOCUS A11128 3400 bp DNA linear PAT 07-DEC-1993  
 DEFINITION Synthetic nucleotide sequence of the leftward reading frame of the Bam B-fragment encoding gp 250/350.  
 ACCESSION A11128  
 VERSION A11128  
 KEYWORDS A11128.1 GI:490970  
 SOURCE synthetic construct  
 ORGANISM synthetic construct  
 artificial sequences.  
 REFERENCE 1 (bases 1 to 3400)  
 Wolf,H.J.  
 DNA sequences of the EBV genome, recombinant DNA molecules,  
 processes for producing EBV-related antigens, diagnostic  
 compositions and pharmaceutical compositions containing said  
 antigens  
 Patent: EP 0173254-A 3 05-MAR-1986;  
 Wolf, Hans Joachim  
 Location/Qualifiers

JOURNAL  
 FEATURES

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/mol_type="genomic DNA"
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556..3279
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/db_xref="GI:490971"
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SOVLPGDKRNTICSGTESHVPSGGLITSTSPVATPIPGGVASISLALTRPVRFLG
NSILVYFVSNGPKASGDYCIQSNIVFSDIIPASQDMPTNTDITVGDNATYSVP
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Query Match
Best Local Similarity 100.0%; Score 20; DB 6; Length 3400;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CCTTAGAGGAGCAAGTCCC 20
Db      2441 CCTTAGAGGAGCAAGTCCC 2460

RESULT 7
LOCUS      E01006      3400 bp      DNA      linear      PAT 29-SEP-1997
DEFINITION      CDNA encoding p350 of Epstein-Barr virus.
ACCESSION      E01006
KEYWORDS      E01006.1 GI:2169265
SOURCE      JP 1986257188-A/2.
ORGANISM      Human herpesvirus 4 (Epstein-Barr virus)
REFERENCE      Human herpesvirus 4
VIRUSES; dsDNA viruses, no RNA stage; Herpesviridae;
Gammaherpesvirinae; Lymphocryptovirus.
1 (bases 1 to 3400)
Hansu, Y.B.
DNA SEQUENCE OF EBV GENOM, RECOMBINED DNA MOLECULE AND EBV RELATED
ANTIGEN AND DIAGNOSTIC COMPOSITION CONTAINING SAID ANTIGEN AND
PRODUCTION OF PREPARATION
Patent: JP 1986257188-A 2 14-NOV-1986;
HANSU YOTSURO BUNORUFU
OS      Epstein-Barr virus
PN      JP 1986257188-A/2
PD      14-NOV-1986
PE      23-AUG-1985 JP 1985185661
PR      23-AUG-1984 EP 84 84110089, 23-AUG-1984 EP 84 84110090 PI
HANSU YOTSURO BUNORUFU
PC      C12N15/00, A61K39/245, C07H21/04, C07K13/00, C12N1/00, G01N33/569,
PC      G01N33/577//
PC      C12P21/02, (C12N1/00, C12R1:19);
CC      strandedness: Double;
CC      topology: Linear;
CC      hypothetical: No;
CC      anti-sense: No;
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/db_xref="GI:490971"
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KMNCKSTLTAVYRAQGLDVTPLSLPTSAQSNSTYKEMGNIDIECIMDEEI
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BASE COUNT      904 a      1060 c      732 g      704 t
ORIGIN

Query Match
Best Local Similarity 100.0%; Score 20; DB 6; Length 3400;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CCTTAGAGGAGCAAGTCCC 20
Db      2441 CCTTAGAGGAGCAAGTCCC 2460

RESULT 8
LOCUS      AR049357      3833 bp      DNA      linear      PAT 29-SEP-1999
DEFINITION      Sequence 18 from patent US 5824508.
ACCESSION      AR049357
KEYWORDS      AR049357.1 GI:6005396
SOURCE      Unknown.
ORGANISM      Unknown.
REFERENCE      Unclassified.
1 (bases 1 to 3833)
AUTHORS      Spaete, R. and Jackman, W.T.
TITLE      Non-splicing variants of gp350/220
JOURNAL      Patent: US 5824508-A 18-20-OCT-1998;
FEATURES      Location/Qualifiers
source      1..3833
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ORIGIN

Query Match
Best Local Similarity 100.0%; Score 20; DB 6; Length 3833;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 CCTTAGAGGAGCAAGTCCC 20
Db      2899 CCTTAGAGGAGCAAGTCCC 2918

RESULT 9
LOCUS      A11178      5019 bp      DNA      linear      PAT 07-DEC-1993
DEFINITION      Synthetic nucleotide sequence for the fusion protein encoded by
PURUP1.9.
ACCESSION      A11178
KEYWORDS      A11178.1 GI:490997
SOURCE      synthetic construct
ORGANISM      synthetic construct
REFERENCE      artificial sequences.
1 (bases 1 to 5019)
WOLF, H.J.
DNA sequences of the EBV genome, recombinant DNA molecules,
processes for producing EBV-related antigens, diagnostic
compositions and pharmaceutical compositions containing said
antigens
Patent: EP 0173254-A 5 05-MAR-1986;
WOLF, Hans Joachim
JOURNAL      Location/Qualifiers
FEATURES      1..5019
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/mol_type="genomic DNA"

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of the P3HR-1 deletion junction and characterization of the NotI repeat units that form part of the template for an abundant 12-O-tetradecanoylphorbol-13-acetate-induced mRNA transcript  
JOURNAL  
MEDLINE  
PUBMED  
83294686  
6310141  
8  
Bankier,A.T., Deininger,P.L., Satchwell,S.C., Baer,R., Farrell,P.J. and Barrell,B.G.  
DNA sequence analysis of the EcoRI DheI fragment of B95-8 Epstein-Barr virus containing the terminal repeat sequences  
Mol. Biol. Med. 1 (4), 425-445 (1983)  
JOURNAL  
MEDLINE  
85060428  
6094955  
9  
Farrell,P.J., Bankier,A., Seguin,C., Deininger,P. and Barrell,B.G.  
Latent and lytic cycle promoters of Epstein-Barr virus  
EMBO J. 2 (8), 1331-1338 (1983)  
JOURNAL  
MEDLINE  
20311131  
10872327  
10  
Jones,M.D., Foster,L., Sheedy,T. and Griffin,B.E.  
The EB virus genome in Daudi Burkitt's lymphoma cells has a deletion similar to that observed in a non-transforming strain (P3HR-1) of the virus  
EMBO J. 3 (4), 813-821 (1984)  
JOURNAL  
MEDLINE  
84207939  
6327290  
11  
Biggin,M., Farrell,P.J. and Barrell,B.G.  
Transcription and DNA sequence of the BamHI I fragment of B95-8 Epstein-Barr virus  
EMBO J. 3 (5), 1083-1090 (1984)  
JOURNAL  
MEDLINE  
84236104  
6203743  
12  
Yates,J., Warren,N., Reisman,D. and Sugden,B.  
A cis-acting element from the Epstein-Barr viral genome that permits stable replication of recombinant plasmids in latently infected cells  
Proc. Natl. Acad. Sci. U.S.A. 81 (12), 3806-3810 (1984)  
JOURNAL  
MEDLINE  
84222045  
6328526  
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Gibson,T., Stockwell,P., Ginsburg,M. and Barrell,B.  
Homology between two EBV early genes and HSV ribonucleotide reductase and 38K genes  
Nucleic Acids Res. 12 (12), 5087-5099 (1984)  
JOURNAL  
MEDLINE  
84247360  
6330697  
14 (bases 1 to 171823)  
Baer,R.J., Bankier,A.T., Biggin,M.D., Deininger,P.L., Farrell,P.J., Gibson,T.J., Hatfull,G.F., Hudson,G.S., Satchwell,S.C., Seguin,C., Tuffnell,P.S. and Barrell,B.G.  
DNA sequence and expression of the B95-8 Epstein-Barr virus genome  
Nature 310 (5974), 207-211 (1984)  
JOURNAL  
MEDLINE  
84270667  
6087149  
15  
Bodasect,M. and Pericandec,M.  
Cluscted alternative splice sites in Epstein-Barr virus RNAs  
Nucleic Acids Res. 15 (14), 5887 (1987)  
JOURNAL  
MEDLINE  
87289053  
3039467  
16  
Lau,G., Pericandec,M. and Farrell,P.J.  
A spliced Epstein-Barr virus gene expressed in immortalized lymphocytes is created by circularization of the linear viral genome  
EMBO J. 7 (3), 769-774 (1988)  
JOURNAL  
MEDLINE  
88283646  
2840285  
REFERENCE

AUTHORS  
TITLE  
JOURNAL  
MEDLINE  
PUBMED  
91021036  
2171209  
18 (bases 1 to 171823)  
Hatfull,G.F., Barrell,B.G., Quinn,J. and McGeoch,D.  
Unpublished  
19  
Binn,U.K., Amon,W. and Farrell,P.J.  
Induction of Epstein-Barr virus late promoters on small plasmids in the EBV late lytic cycle requires ori L $\gamma$   
Unpublished  
20 (bases 1 to 171823)  
Farrell,P.J.  
Direct Submission  
Submitted (01-AUG-2002) Farrell P., Ludwig Institute for Cancer Research, Imperial College School of Medicine, St. Mary's Campus, Norfolk Place London W2 1PG  
Construction:  
This sequence was assembled from B95-8 EBV [14] and Raji EBV [18] with sequence corrections [16, 19]. The number of major internal repeat units has been reduced from 11.6 [14] to a more typical 7.6 and the B95-8 deletion sequences have been restored to give a sequence more representative of wild type EBV.  
Numbering  
like the modified B95-8 sequence [14, 16] accession number V01555, this sequence starts 1 base to the left of the EcoRI site separating EcoRI DheI from EcoRI I (ie the first A of AGAATTC).  
Location/Qualifiers  
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/mol\_type="genomic DNA"  
/strain="B95-8"  
/db\_xref="taxon:10376"  
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/mol\_type="genomic DNA"  
/strain="B95-8"  
/db\_xref="taxon:10376"  
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/mol\_type="genomic DNA"  
/strain="Raji"  
/db\_xref="taxon:10376"  
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join(166103..166458,58..272,360..458,540..768,871..951,1026..1196,1280..1495,1574..1680)  
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PRDSSQHTYERAGKSNPVCPLPYVAPYFLWLAIAASCTFASVSYVYATGLAS  
LILAAVASSYAAQOKLTPVTVAVTFAICTWETIDPDPNSLTFALAAAG  
LOGIYLVMLVLLIAYRRMRRLVCGIMFLACVLIVAVQLSPILGAVVVS  
MTLILAVLWVSSPGAGTGAALITLAAALALASLITGLNTTMTLMLTLV  
VLICSSGSCPLSKILARLEFLVALALLALASALIAAGSILOTNFKSLSTFEIPL  
RMLLLIVAGLIFLITLITMGSGNRTGAPVMCIGILTWAGAVMLTVMSNTLISA  
WILTRGFLIFLIGPALFGVIRRCRYCCYCLTLSEBRPPTRYRTV"  
58..272  
/gene="LMP2"  
/number=2  
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/gene="LMP2"  
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/product="terminal protein LMP2B"  
/protein\_id="CAD53383.1"

Query Match	100.0%;	Score 20;	DB 14;	Length 171823;
Best Local Similarity	100.0%;	Pred. No. 0.55;		
Matches	20;	Conservative	0;	Mismatches 0;
			Indels	0;
			Gaps	0;

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QY      1 CCTTAGGAGAACAGTCCC 20
          |||||
Db      77980 CCTTAGGAGGAACAAGTCCC 77961

```

LOCUS	EBV	172281 bp	DNA	circular	VRL 20-SEP-1999
DEFINITION	Epstein-Barr virus (EBV) genome, strain B95-8.				
ACCESSION	V01555	J02070	K01729	K01730	V01554 X00498 X00499 X00784
VERSION	V01555.1	GI:59074			
KEYWORDS	DNA polymerase; EBNA; genome; ribonucleotide reductase; tandem repeat; terminal repeat.				
SOURCE	Human herpesvirus 4 (Epstein-Barr virus)				
ORGANISM	Human herpesvirus 4				

REFERENCE	8 (bases 159853 to 172281)
AUTHORS	Bankier,A.T., Deininger,P.L., Satchell,S.C., Baer,R., Farrell,P.J. and Barrell,B.G.
TITLE	DNA sequence analysis of the EcoRI DheI fragment of B95-8 Epstein-Barr virus containing the terminal repeat sequences
JOURNAL	Mol. Biol. Med. 1 (4), 425-445 (1983)
MEDLINE	85060428
PUBMED	6094955
REFERENCE	9 (bases 1 to 172281)
AUTHORS	Farrell,P.J., Bankier,A., Seguin,C., Deininger,P. and Barrell,B.G.
TITLE	Latent and lytic cycle promoters of Epstein-Barr virus
JOURNAL	EMBO J. 2 (8), 1331-1338 (1983)
MEDLINE	20331131
PUBMED	10872327
REFERENCE	10 (bases 45415 to 52824)
AUTHORS	Jones,M.D., Foster,L., Sheedy,T. and Griffin,B.E.
TITLE	The EB virus genome in Daudi Burkitt's lymphoma cells has a

REFERENCE  
AUTHORS  
TITLE  
JOURNAL

## COMMENT

19 (bases 1 to 172281)  
 Farrell, P.J.  
 Direct Submission  
 Submitted (18-MAR-1988) Farrell P., Ludwig Institute for Cancer  
 Research, St. Mary's Hospital Medical School, Norfolk Place London  
 W2 1PG

## CDS

Listed under this feature are all known protein coding regions as well as all the major open reading frames in the sequence. In general the term major is taken as the longest frame in a particular region taking into account the adjacent longest frames and likely transcription signals. Note that on this basis some long overlapping frames have been excluded and on the other hand some small frames have been included which might represent exons or genes because they occur in a logical combination with other features or because of some other experimental data. The reading frames are named according to the Bam HI fragment in which they start. eg BAF3 is the third leftward frame starting in Bam HI fragment A. BOKFI is the first rightward frame in Bam HI fragment O. If there is an obvious PATA sequence followed by an in frame Met codon that satisfies the rules of Kozak [12] in that there is a purine at -3 and/or a G at +4 then the reading frame is numbered from the A of the ATG to the base preceding the termination codon. If there is no obvious initiation codon or there is a substantial reading frame in phase before the ATG then the reading frame is numbered from the first base of the first codon.

## SITES OF POLYA SIGNALS

This feature lists all occurrences of the sequence AATAA which is found normally approximately 20 bases upstream of the mRNA processing/polyA addition site. The rarely used homolog AATAA is only listed when it is found in a position close to the end of a major reading frame.

## SITES OF DONOR AND ACCEPT SEQUENCES

This is not a comprehensive listing of all such sequences and only the positions of a few have been noted because they occur in potentially interesting positions. The number quoted in the table is the position of the terminal base in the intron in each case. Restriction enzyme SITES.

Only the positions of the sites Bam HI (BAM) are listed.

## RPT

This feature is used to define repetitive sequences.

## SITE DEL.

This feature defines deletions in B95-8 with respect to other strains such as Raji and also to deletions in other strains such as BJH1 and DAUDI with respect to B95-8.

## SITE HPN

Denotes sequences with twofold symmetry ie could form hairpin loops. This is not a comprehensive list - only a few occurrences noted.

## ORGRPL

Denotes the region that encompasses an origin of replication (ori p). [13].

## NUMBERING

The DNA sequence of B95-8 EBV has been revised [19]. The original (Baer et al, 1984) base 359 has been deleted so the new sequence around that position reads TCAGCTTT. To avoid renumbering the entire sequence, position 1 has been moved 1 base to the left of the EcoRI site separating EcoRI Dnet from EcoRI I (ie the first A of AGAATTC).

## FEATURES

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1.172281

/organism="Human herpesvirus 4"

/mol\_type="genomic DNA"

/strain="B95-8"

/db\_xref="taxon:10376"

mrna 58..272

Query Match 100.0%; Score 20; DB 14; Length 172281;

Best Local Similarity 100.0%; Pred. No. 0.55; Indels 0; Gaps 0;

Matches 20; Conservative 0; Mismatches 0

1 CCTAGAGAGAACAGTCCC 20

9

DB 90268 CCTAGAGAGAACAGTCCC 90249

## RESULT.15

## LOCUS

HSAB958RAJ/c 184113 bp DNA linear

## DEFINITION

Epstein-Barr virus, artifactual joining of B95-8 complete genome and the sequences from Raji of the large deletion found in B95-8.

Accession M80517.1 GI:330330

## VERSION

## KEYWORDS

## SOURCE

## ORGANISM

Human herpesvirus 4 (Epstein-Barr virus)

Viruses; dsDNA viruses, no RNA stage; Herpesviridae; Gammaherpesvirinae; Lymphocryptovirus.

## REFERENCE

## AUTHORS

Baer, R.J., Bankier, A.T., Biggin, M.D., Deininger, P.L., Farrell, P.J., Gibson, T.J., Hatfull, G.F., Hudson, G.S., Satchwell, S.C., Segun, C., Tufnell, P.S. and Barrell, B.G.

DNA sequence and expression of the B95-8 Epstein-Barr virus genome

Nature 310 (5974), 207-211 (1984)

## MEDLINE

84270667

## PUBMED

6087149

## REFERENCE

## AUTHORS

Parker, B.D., Bankier, A., Satchwell, S., Barrell, B. and Farrell, P.J.

Sequence and transcription of Raji Epstein-Barr virus DNA spanning

the B95-8 deletion region

Virology 179 (1), 339-346 (1990)

## MEDLINE

91021036

## PUBMED

2171209

## REFERENCE

## AUTHORS

Sample, J., Brooks, L., Sample, C., Young, L., Rowe, M., Gregory, C.,

Rickinson, A. and Kieff, E.

Restricted Epstein-Barr virus protein expression in Burkitt

lymphoma is due to a different Epstein-Barr nuclear antigen 1

transcriptional initiation site

Proc. Natl. Acad. Sci. U.S.A. 88 (14), 6343-6347 (1991)

## MEDLINE

91296817

## PUBMED

1648738

## REFERENCE

## AUTHORS

Jensen, H.B.

GenBank Curator Program

Unpublished (1992)

## JOURNAL

## COMMENT

Original source text: Human herpesvirus 4 DNA. The B95-8 genome (V01555) has a large deletion in the right side of the genome which has been sequenced in Raji (M35547). These sequences have been joined to form an extended and more complete, although artifactual, EBV sequence.

For features, refer to feature tables of V01555 and M35547.

## FEATURES

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/mol\_type="genomic DNA"

/db\_xref="taxon:10376"

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152009..152012

/note="Overlap of B95-8 and Raji sequences at B95-8 deletion point (corresponds to 152,009-152,012 in V01555, and 1-4 in M35547)"

153013..163839

/note="Raji sequences (corresponds to 5-11,831 of M35547)"

163840..163843

/note="Overlap of B95-8 and Raji sequences at B95-8 deletion point (corresponds to 152,009-152,012 of V01555, and 11,832-11,835 of M35547)"

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/note="B95-8 sequences (corresponds to 152,013-172,282 of V01555)"

## misc\_feature

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## ORIGIN

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 9c268 CCTTAGAGGACAGTCCC 90249

RESULT 16  
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 LOCUS Rattus norvegicus clone CH230-522E23, \*\*\* SEQUENCING IN PROGRESS  
 DEFINITION  
 AC142486  
 AC142486.2 GI:29423830  
 HTG; HTGS-PHASEL  
 Rattus norvegicus (Norway rat)  
 Rattus norvegicus  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;  
 Rattus.

REFERENCE  
 1 (bases 1 to 163273)  
 Muzny, D., Marre, M., Metzker, M., Lee, A., Abramson, S., Adams, C., Alder, J.,  
 Allen, C., Allen, H., Alsbrooks, S., Amin, A., Anguiano, D.,  
 Anyalebechi, V., Aoyagi, A., Ayodeji, M., Baca, E., Baden, H.,  
 Baldwin, D., Bandaranaike, D., Barber, M., Barnstead, M., Benahmed, F.,  
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 Bryant, N., Buhay, C., Burch, P., Burrell, K., Calderon, E.,  
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 Chacko, J., Chavez, D., Chen, G., Chen, Y., Chen, Z., Chu, J.,  
 Cleveland, C., Cockrell, R., Cox, C., Coyle, M., Cree, A., D'Souza, L.,  
 Davila, M. L., Davis, C., Davy-Carroll, L., De Anda, C., Dederich, D.,  
 Delgado, O., DeGron, S., Deramo, C., Ding, Y., Dinh, H., Divya, K.,  
 Draper, H., Dugan-Rocha, S., Dunn, A., Durbin, K., Duval, B., Eaves, K.,  
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 Fraser, C. M., Gabisi, A., Ganta, R., Garcia, A., Garner, T., Gervais, M.,  
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 Jackson, L., Jacob, L., Jiang, H., Johnson, B., Johnson, R., Jolivet, A.,  
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 Yen, J., Yoon, L., Yoon, V., Yu, F., Zhang, J., Zhou, J., Zhou, X.,  
 Zhao, S., Dunn, D., von Niederhausern, A., Weiss, R., Smith, D. R.,  
 Holt, R. A., Smith, H. O., Weinstock, G. and Gibbs, R. A.

TITLE  
 Direct Submission  
 Unpublished

REFERENCE  
 2 (bases 1 to 163273)  
 Worley, K.C.  
 Direct Submission  
 Submitted (31-MAR-2003) Human Genome Sequencing Center, Department  
 of Molecular and Human Genetics, Baylor College of Medicine, One  
 Baylor Plaza, Houston, TX 77030, USA  
 3 (bases 1 to 163273)  
 Worley, K.C.  
 Direct Submission  
 Submitted (01-APR-2003) Human Genome Sequencing Center, Department  
 of Molecular and Human Genetics, Baylor College of Medicine, One  
 Baylor Plaza, Houston, TX 77030, USA  
 On Apr 1, 2003 this sequence version replaced gi:29374196.

COMMENT  
 Center: Baylor College of Medicine  
 Center code: BCM  
 Web site: <http://www.hgsc.bcm.tmc.edu/>  
 Contact: hgsc-help@bcm.tmc.edu  
 Project Information  
 Center project name: KEMO  
 Center clone name: CH230-522E23  
 Summary Statistics  
 Sequencing vector: plasmid  
 Chemistry: Dye-terminator Big Dye 100% of reads  
 Assembly program: Phrap, version 0.990329  
 Consensus quality: 140865 bases at least Q40  
 Consensus quality: 147625 bases at least Q30  
 Consensus quality: 152515 bases at least Q20  
 Estimated insert size: 149404; sum-of-contigs estimation  
 Quality coverage: 2x in Q20 bases; sum-of-contigs estimation

NOTE: Estimated insert size may differ from sequence length  
 (see [http://www.hgsc.bcm.tmc.edu/docs/Genbank\\_draft\\_data.html](http://www.hgsc.bcm.tmc.edu/docs/Genbank_draft_data.html)).  
 NOTE: This is a 'working draft' sequence. It currently  
 consists of 49 contigs. The true order of the pieces  
 is not known and their order in this sequence record is  
 arbitrary. Gaps between the contigs are represented as  
 runs of N, but the exact sizes of the gaps are unknown.  
 This record will be updated with the finished sequence  
 as soon as it is available and the accession number will  
 be preserved.

1 1102: contig of 1102 bp in length  
 \* 1103 1202: gap of unknown length  
 \* 1203 2433: contig of 1231 bp in length  
 \* 2434 2533: gap of unknown length  
 \* 2534 3667: contig of 1134 bp in length  
 \* 3668 3767: gap of unknown length  
 \* 3768 4781: contig of 1014 bp in length  
 \* 4782 4881: gap of unknown length  
 \* 4882 6332: contig of 1451 bp in length  
 \* 6333 6432: gap of unknown length  
 \* 6433 7726: contig of 1294 bp in length  
 \* 7727 7826: gap of unknown length  
 \* 7827 9239: contig of 1413 bp in length  
 \* 9240 9339: gap of unknown length  
 \* 9340 10646: contig of 1307 bp in length  
 \* 10647 10746: gap of unknown length  
 \* 10747 13007: contig of 2261 bp in length  
 \* 13008 13107: gap of unknown length  
 \* 13108 14650: contig of 1543 bp in length  
 \* 14651 14750: gap of unknown length  
 \* 14751 16039: contig of 1289 bp in length  
 \* 16040 16139: gap of unknown length  
 \* 16140 18332: contig of 2093 bp in length  
 \* 18333 18332: gap of unknown length  
 \* 18333 19866: contig of 1534 bp in length  
 \* 19867 19966: gap of unknown length  
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 \* 21079 21178: gap of unknown length  
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 \* 22966 23065: gap of unknown length  
 \* 23066 25141: contig of 2076 bp in length  
 \* 25142 25241: gap of unknown length

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*	101853	108908:	contig of 7056 bp in length
*	108909	109008:	gap of unknown length
*	109009	115481:	contig of 6473 bp in length
*	115482	115581:	gap of unknown length
*	115582	122306:	contig of 6725 bp in length
*	122307	122406:	gap of unknown length
*	122407	130001:	contig of 7595 bp in length
*	130002	130101:	gap of unknown length
*	130102	138191:	contig of 8096 bp in length
*	138192	138291:	gap of unknown length
*	138292	150672:	contig of 12361 bp in length
*	150673	150772:	gap of unknown length
*	150773	163273:	contig of 12501 bp in length.

Query Match 85.0% Score 17; DB 2; Length 163273;

Best Local Similarity 100.0% Pared No. 22; Mismatches 0; Indels 0; Gaps 0;

DB 66325 CTTAGAGGAGAACAGTC 66309

RESULT 17  
AC113548/c  
LOCUS  
DEFINITION  
MUS MUSCULUS clone RP23-268F15, WORKING DRAFT SEQUENCE, 11  
unordered pieces.  
AC113548  
AC113548.5 GI:28913170  
HTG, HTGS, PHASE1, HTGS, DRAFT.  
MUS MUSCULUS (house mouse)  
ORGANISM  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.  
REFERENCE  
Birren, B., Nussbaum, C. and Lander, E.  
Mus musculus, clone RP23-268F15  
Unpublished  
2 (bases 1 to 193466)

Birren, B., Linton, L., Nussbaum, C., Lander, E., All, A., Allen, N.,  
Anderson, S., Barna, N., Bastien, V., Boguslavsky, L., Boukhalter, B.,  
Brown, A., Camarata, J., Campolano, A., Chang, J., Chazaro, B.,  
Choepel, Y., Colangelo, M., Collins, S., Collymore, A., Cook, A.,  
Cooke, P., Dearellano, K., Dewar, K., Diaz, J.S., Dodge, S., Faro, S.,  
Ferreira, P., Fitzhugh, W., Gage, D., Galagan, J., Gardyna, S.,  
Ginde, S., Gord, S., Goyette, M., Graham, L., Grand-Pierre, N.,  
Hagos, B., Horton, L., Hulme, W., Iliev, I., Johnson, R., Jones, C.,  
Kamat, A., Karatas, A., Kells, C., Laroque, K., Lamazares, R.,  
Landre, T., Lehoczy, J., Levine, R., Liu, G., Maclean, C.,  
Macdonald, P., Major, J., Marquis, N., Matthews, C., McCarthy, M.,  
McEwan, P., McKernan, K., Meldrum, J., Meneus, L., Mihova, T.,  
Mlenga, V., Murphy, T., Naylor, J., Nguyen, C., Nicol, R., Norbu, C.,  
Norman, C.H., O'Connor, T., O'Donnell, P., O'Neill, D., Oliver, J.,  
Peterson, K., Phunkhang, P., Pierre, N., Pollara, V., Raymond, C.,  
Retta, R., Rieback, M., Riley, R., Rise, C., Rogov, P., Roman, J.,  
Roselli, M., Roy, A., Santos, R., Schauer, S., Schupack, R., Seaman, S.,  
Severy, P., Spencer, B., Stange-Thomann, N., Stojanovic, N.,  
Strauss, K., Subramanian, A., Talamas, J., Testfaye, S., Theodore, J.,  
Topham, K., Travers, M., Travis, N., Trigilio, J., Vassiliev, H.,  
Viel, R., Vo, A., Wilson, B., Wu, X., Wymann, D., Ye, W.J., Young, G.,  
Zainoun, J., Zembek, L., Zimmer, A. and Zody, M.

TITLE  
JOURNAL  
Direct Submission  
Submitted (01-MAR-2002) Whitehead Institute/MIT Center for Genome  
Research, 320 Charles Street, Cambridge, MA 02141, USA  
3 (bases 1 to 193466)

REFERENCE  
AUTHORS  
Birren, B., Nussbaum, C., Lander, E., Abouelleil, A., Allen, N.,  
Anderson, S., Arachchi, H.M., Barna, N., Bastien, V., Bloom, T.,  
Boguslavsky, L., Boukhalter, B., Camarata, J., Chang, J., Choepel, Y.,  
Collymore, A., Cook, A., Cooke, P., Corum, B., Dearellano, K.,  
Diaz, J.S., Dodge, S., Dooley, K., Dorris, L., Erickson, J., Faro, S.,  
Ferreira, P., Fitzgerald, M., Gage, D., Galagan, J., Gardyna, S.,  
Graham, L., Grand-Pierre, N., Hateg, N., Hasopian, D., Hagos, B.,  
Hall, J., Horton, L., Hulme, W., Iliev, I., Johnson, R., Jones, C.,  
Kamat, A., Karatas, A., Kells, C., Landers, T., Levine, R.,  
Lindblad-Toh, K., Liu, G., Lui, A., Mabbitt, R., Maclean, C.,  
Macdonald, P., Major, J., Manning, J., Matthews, C., McCarthy, M.,  
Meldrum, J., Meneus, L., Mihova, T., Mlenga, V., Murphy, T., Naylor, J.,  
Nguyen, C., Nicol, R., Norbu, C., O'Connor, T., O'Donnell, P.,  
O'Neill, D., Oliver, J., Peterson, K., Phunkhang, P., Pierre, N.,  
Rachupka, A., Ramsamy, U., Raymond, C., Retta, R., Rise, C., Rogov, P.,  
Roman, J., Schauer, S., Schupack, R., Seaman, S., Severy, P., Smith, C.,  
Spencer, B., Stange-Thomann, N., Stojanovic, N., Stubbs, M.,  
Talamas, J., Testfaye, S., Theodore, J., Topham, K., Travers, M.,  
Vassiliev, H., Venkataraman, V.S., Viel, R., Vo, A., Wilson, B., Wu, X.,  
Wymann, D., Young, G., Zainoun, J., Zembek, L., Zimmer, A. and Zody, M.

TITLE  
JOURNAL  
Direct Submission  
Submitted (11-MAR-2003) Whitehead Institute/MIT Center for Genome  
Research, 320 Charles Street, Cambridge, MA 02141, USA  
On Mar 11, 2003 this sequence version replaced gi:28626751.  
All repeats were identified using RepeatMasker:  
Smit, A.F.A. & Green, P. (1996-1997).  
http://ftp.genome.washington.edu/BM/RepeatMasker.html  
Genome Center



Center: Whitehead Institute/ MIT Center for Genome Research  
 Center code: WIBR  
 Web site: <http://www-seq.wi.mit.edu>  
 Contact: [sequence.submissions@genome.wi.mit.edu](mailto:sequence.submissions@genome.wi.mit.edu)

----- Project Information -----  
 Center project name: L24242  
 Center clone name: 268\_F\_15

----- Summary Statistics -----  
 Sequencing vector: plasmid; n/a; 100% of reads  
 Chemistry: Dye-terminator Big Dye; 100% of reads  
 Assembly program: Phrap; version 0.960731  
 Consensus quality: 191143 bases at least Q40  
 Consensus quality: 191818 bases at least Q30  
 Consensus quality: 192162 bases at least Q20  
 Insert size: 192000; agarose-fp  
 Insert size: 192466; sum-of-coverage  
 Quality coverage: 8.5 in Q20 bases; agarose-fp  
 Quality coverage: 8.5 in Q20 bases; sum-of-coverage

----- NOTE: This is a 'working draft' sequence. It currently consists of 11 contigs. The true order of the pieces is not known and their order in this sequence record is arbitrary. Gaps between the contigs are represented as runs of N, but the exact sizes of the gaps are unknown. This record will be updated with the finished sequence as soon as it is available and the accession number will be preserved. -----

1 4933: contig of 4933 bp in length  
 4934 5033: gap of 100 bp  
 5034 6097: contig of 1064 bp in length  
 6098 6197: gap of 100 bp  
 6198 7378: contig of 1181 bp in length  
 7379 7478: gap of 100 bp  
 7479 9778: contig of 2300 bp in length  
 9779 9878: gap of 100 bp  
 9879 11949: contig of 2071 bp in length  
 11950 12049: gap of 100 bp  
 12050 15641: contig of 3592 bp in length  
 15642 15741: gap of 100 bp  
 15742 19456: contig of 3715 bp in length  
 19457 39905: contig of 20349 bp in length  
 39906 40005: gap of 100 bp  
 40006 67129: contig of 27124 bp in length  
 67130 67230: gap of 100 bp  
 67231 104769: contig of 37540 bp in length  
 104770 104870: gap of 100 bp  
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 vector\_side:right"

BASE COUNT 53362 a 45171 c 42934 g 50990 t 1009 others  
 ORIGIN

Query Match 85.0%; Score 17; DB 2; Length 193466;  
 Best Local Similarity 100.0%; Pred. No. 21;  
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 TAGGAGACAAAGTCCC 20  
 Db 147268 TAGGAGACAAAGTCCC 147252

RESULT 18  
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 DEFINITION human SRS A005R48, sequence tagged site.  
 ACCESSION G20457.1 GI:1340794  
 VERSION G20457.1 GI:1340794  
 KEYWORDS SRS; SRS sequence; primer; sequence tagged site.  
 SOURCE Homo sapiens (human)  
 ORGANISM Homo sapiens  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
 1 (bases 1 to 204)  
 Adams, M.D.  
 JOURNAL Unpublished (1996)  
 COMMENT

REFERENCE  
 AUTHORS  
 JOURNAL  
 COMMENT

Contact: Mark Adams  
 The Institute for Genomic Research  
 9712 Medical Center Dr., Rockville, MD 20850  
 Email: [mdadams@tigr.org](mailto:mdadams@tigr.org)

Primer A: GTATCTCTTAGGACAAAGTA  
 Primer B: ATTGAGTCTTAGAATTACTTA  
 SRS size: 204  
 PCR Profile:  
 Denaturation: 96C 5min  
 Anneal: 54C 30sec  
 Extend: 72C 30sec  
 Denature: 95C 30sec  
 FinalExtend: 72C 5min  
 Cycles: 30

Protocol:  
 GenomicDNA: 25 ng  
 Primer: 0.43 uM each  
 dNTPs: 230 uM each  
 AmpliTaq: 0.5 units  
 Tagstart Ab: 0.5 units  
 Total Volume: 10 uL  
 Cycles: 30

Buffer:  
 Tris-HCl pH8.8: 100 mM  
 KCl: 500 mM  
 MgCl2: 20 mM  
 Triton X-100: 1%  
 Concentration: 10X

Prepared with primer pairs derived from THCI05423: GenBank  
 Accession Numbers: H80345, H70110, F11453, F13120, R24326, R24340,  
 R24852, T75305, Z42603.  
 Location/Qualifiers  
 1. 204  
 /organism="Homo sapiens"  
 /mol\_type="genomic DNA"  
 /db\_xref="taxon:9606"

FEATURES  
 source



STS  
primer\_bind 1. .204  
primer\_bind 1. .23  
BASE COUNT 76 a 39 c 34 g 55 t  
ORIGIN

Query Match 80.0%; Score 16; DB 11; Length 204;  
Best Local Similarity 100.0%; Pred. No. 3.1e+02;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 CTTAGGAGGACAGT 17  
|||||  
Db 7 CTTAGGAGGACAGT 22

RESULT 19  
AY079380 1555 bp mRNA linear PLN 18-SEP-2002  
LOCUS Arabidopsis thaliana unknown protein (At2g44020) mRNA, complete  
DEFINITION cds.

ACCESSION AY079380 GI:19310760  
KEYWORDS FLI CDNA.  
SOURCE Arabidopsis thaliana (thale cress)  
ORGANISM Arabidopsis thaliana  
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;  
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;  
rosids; eustosids II; Brassicales; Brassicaceae; Arabidopsis.

REFERENCE 1 (bases 1 to 1555)  
Yamada, K., Liu, S.X., Sakano, H., Pham, P.K., Banh, J., Egu, P.,  
Lee, J.M., Toriumi, M., Yu, G., Brooks, S., Chao, Q., Chen, H.,  
Karlin-Neumann, G., Kim, C., Lam, B., Miranda, M., Nguyen, M.,  
Palm, C.J., Shinn, P., Southwick, A., Davis, R.W., Ecker, J.R. and  
Theologis, A.

TITLE Arabidopsis Open Reading Frame (ORF) Clones  
JOURNAL Unpublished  
REFERENCE 2 (bases 1 to 1555)

Yamada, K., Banh, J., Chan, M.M., Chang, C.H., Chang, E., Dale, J.M.,  
Deng, J.M., Goldsmith, A.D., Lee, J.M., Onodera, C.S., Quach, H.L.,  
Tang, C.C., Toriumi, M., Wu, H.C., Yamamura, Y., Yu, G., Bowser, L.,  
Carninci, P., Chen, H., Cheuk, R., Hayashizaki, Y., Ishida, J.,  
Jones, T., Kamiya, A., Karlin-Neumann, G., Kawai, J., Kim, C., Lam, B.,  
Lin, J., Meyers, M.C., Miranda, M., Narusaka, M., Nguyen, M., Palm, C.J.,  
Sakurai, T., Satou, M., Seki, M., Shinn, P., Southwick, A.,  
Shinozaki, K., Davis, R.W., Ecker, J.R. and Theologis, A.

TITLE Direct Submission  
JOURNAL Submitted (19-FEB-2002) Plant Gene Expression Center, 800 Buchanan  
Street, Albany, CA 94710, USA

COMMENT The RIKEN Genomic Sciences Center (GSC) members carried out the  
collection and clustering of RAFL cDNAs (RAFL cDNA : RIKEN  
Arabidopsis Full-length cDNA) : Seki, M., Narusaka, M., Ishida, J.,  
Satou, M., Kamiya, A., Sakurai, T., Carninci, P., Kawai, J.,  
Hayashizaki, Y. and Shinozaki, K.

The Salik, Stanford, PGEC (SSP) Consortium members constructed and  
sequenced the pUNI (ORF) clones using the RAFL cDNAs: Yamada, K.,  
Banh, J., Chan, M.M., Chang, C.H., Chang, E., Dale, J.M., Deng, J.M.,  
Goldsmith, A.D., Lee, J.M., Onodera, C.S., Quach, H.L., Tang, C.C.,  
Toriumi, M., Wu, H.C., Yamamura, Y., Yu, G., Bowser, L., Chen, H.,  
Cheuk, R., Jones, T., Karlin-Neumann, G., Kim, C., Lam, B., Lin, J.,  
Meyers, M.C., Miranda, M., Nguyen, M., Palm, C.J., Shinn, P.,  
Southwick, A., Davis, R.W., Ecker, J.R. and Theologis, A.

Yamada, K. (SSP/PGEC) and Seki, M. (RIKEN GSC) contributed equally  
to this work. Shinozaki, K. (RIKEN GSC) and Theologis, A. (SSP  
/PGSC) contributed equally to this work as PIs.

Annotation is based on the January 2002 version of the Arabidopsis  
genome submitted to GenBank.

FEATURES  
source 1. .1555  
Location/Qualifiers  
/organism="Arabidopsis thaliana"  
/mol\_type="mRNA"

gene  
CDS

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SVAYLVIGVSPRDIGPMVQYPIILGKRVGTMIPLDYLLISIGPKKIVARMEKR  
SYIVGNLEETVKNPVDCLISFGVKELLPILIAQVQILGLPVAKNSIQYFSLK  
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ELMKNSYFYKTEKMRPKMELVEYEPFTYSLSRIKPRYQGLQSGKTSINMPLNC  
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BASE COUNT 463 a 271 c 372 g 449 t

Query Match 80.0%; Score 16; DB 8; Length 1555;  
Best Local Similarity 100.0%; Pred. No. 2e+02;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 CTTAGGAGGACAGT 17  
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Db 13 CTTAGGAGGACAGT 28

RESULT 20  
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LOCUS Arabidopsis thaliana unknown protein (At2g44020) mRNA, complete  
DEFINITION cds.

ACCESSION AY039920 GI:14532591  
KEYWORDS FLI CDNA.  
SOURCE Arabidopsis thaliana (thale cress)  
ORGANISM Arabidopsis thaliana  
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;  
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;  
rosids; eustosids II; Brassicales; Brassicaceae; Arabidopsis.

REFERENCE 1 (bases 1 to 1767)  
Yamada, K., Liu, S.X., Sakano, H., Pham, P.K., Banh, J., Chung, M.K.,  
Goldsmith, A.D., Lee, J.M., Quach, H.L., Toriumi, M., Yu, G., Bowser, L.,  
Carninci, P., Chen, H., Cheuk, R., Hayashizaki, Y., Ishida, J.,  
Jones, T., Kamiya, A., Karlin-Neumann, G., Kawai, J., Kim, C., Lam, B.,  
Lin, J., Miranda, M., Narusaka, M., Nguyen, M., Palm, C.J., Sakurai, T.,  
Satou, M., Seki, M., Shinn, P., Southwick, A., Shinozaki, K.,  
Davis, R.W., Ecker, J.R. and Theologis, A.

TITLE Arabidopsis Full length cDNA Clones  
JOURNAL Unpublished  
REFERENCE 2 (bases 1 to 1767)

Yamada, K., Liu, S.X., Sakano, H., Pham, P.K., Banh, J., Chung, M.K.,  
Dale, J.M., Gibson, H.A., Goldsmith, A.D., Jiang, P.X., Lee, J.M.,  
Quach, H.L., Tang, C.C., Toriumi, M., Yu, G., Bowser, L., Carninci, P.,  
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Karlin-Neumann, G., Kawai, J., Kim, C., Koesema, E., Lam, B., Lin, J.,  
Meyers, M.C., Miranda, M., Narusaka, M., Nguyen, M., Palm, C.J.,  
Sakurai, T., Satou, M., Seki, M., Shinn, P., Southwick, A., Tracy, S.E.,  
Shinozaki, K., Davis, R.W., Ecker, J.R. and Theologis, A.

Direct Submission  
JOURNAL Submitted (07-JUN-2001) Plant Gene Expression Center, 800 Buchanan  
Street, Albany, CA 94710, USA

COMMENT The RIKEN Genomic Sciences Center (GSC) members carried out the

collection and clustering of RAFL cDNAs (RAFL cDNA : 'RIKEN Arabidopsis Full-length cDNA') : Seki,M., Narusaka,M., Ishida,J., Satou,M., Kamiya,A., Sakurai,T., Carninci,P., Kawai,J., Hayashizaki,Y. and Shinozaki,K.

The Salk, Stanford, PEGC (SSP) Consortium members carried out the sequencing and annotation of the RAFL cDNAs: Yamada,K., Liu,S.X., Sakano,H., Pham,P.K., Banh,J., Chung,M.K., Dale,J.M., Gibson,H.A., Goldsmith,A.D., Jiang,P.X., Lee,J.M., Quach,H.L., Tang,C.C., Toriumi,M., Yu,G., Bowser,L., Chen,H., Cheuk,R., Jones,T., Karlin-Neumann,G., Kim,C., Koeseema,E., Lam,B., Lin,J., Meyers,M.C., Miranda,M., Nguyen,M., Palm,C.J., Shim,P., Southwick,A., Tracy,S.E., Davis,R.W., Ecker,J.R. and Theologis,A.

Yamada,K. (SSP/PEGC) and Seki,M. (RIKEN GSC) contributed equally to this work. Shinozaki,K. (RIKEN GSC) and Theologis,A. (SSP/PEGC) contributed equally to this work as PIs.

Annotation is based on the January 2002 version of the Arabidopsis genome submitted to GenBank.

FEATURES  
source

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ELMKNSYFYKTEMGRPMKELVLPPEFTYLSRSRIKPRYQKQSGIRSSLMPLNC  
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BASE COUNT 520 a 316 c 404 g 527 t  
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Query Match 80.0%; Score 16; DB 8; Length 1767;  
Best local Similarity 100.0%; Pred. No. 1.9e+02;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 CTTAGAGGAAACAGT 17  
|||||  
Db 138 CTTAGAGGAAACAGT 153

RESULT 21  
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LOCUS AX151433  
DEFINITION Sequence 38 from Patent WO0138351.  
ACCESSION AX151433  
VERSION AX151433.1 GI:14533498  
KEYWORDS  
SOURCE Shrimp white spot syndrome virus  
ORGANISM Shrimp white spot syndrome virus

REFERENCE  
AUTHORS Xu,X., Yang,F., He,J., Pham,L.Z., He,M., Ye,Y., Shen,Y. and Kodira,C.  
TITLE Nucleotide sequence of the shrimp white spot syndrome bacilliform virus (wsbv), systems containing this sequence and detection kits  
JOURNAL Patent: WO 0138351-A 38 31-MAY-2001;  
PE Corporation (NY) (US) ; The Third Institute of Oceanography, State Oceanic Administration (CN) ; Sinogenomax Co., Ltd. (CN)

FEATURES  
source

1.2376  
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/mol\_type="genomic DNA"  
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BASE COUNT 787 a 467 c 526 g 596 t  
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Best local Similarity 100.0%; Pred. No. 1.8e+02;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 AGGAGGAAACAGTCCC 20  
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Db 793 AGGAGGAAACAGTCCC 808

RESULT 22  
LOCUS AB002380 6203 bp mRNA linear PRI 06-OCT-2001  
DEFINITION Human mRNA for KIAA0382 gene, partial cds.  
ACCESSION AB002380  
VERSION AB002380.1 GI:2224704  
KEYWORDS KIAA0382.  
SOURCE Homo sapiens (human)  
ORGANISM Homo sapiens  
REFERENCE  
AUTHORS Nagase,T., Ishikawa,K., Nakajima,D., Ohira,M., Seki,N., Miyajima,N., Tanaka,A., Kotani,H., Nomura,N. and Ohara,O.  
TITLE Prediction of the coding sequences of unidentified human genes. VII. The complete sequences of 100 new cDNA clones from brain which can code for large proteins in vitro  
JOURNAL DNA Res. 4 (2), 141-150 (1997)  
MEDLINE 97349984  
PUBMED 9205841  
REFERENCE  
AUTHORS 2 (bases 1 to 6203)  
Ohara,O., Nagase,T., Kikuno,R. and Nomura,N.  
TITLE Direct Submision  
JOURNAL Submitted (28-MAR-1997) Osamu Ohara, Kazusa DNA Research Institute; 1532-3, Yana, Kisarazu, Chiba 292-0812, Japan  
(E-mail:cdnainfo@kazusa.or.jp, Tel:+81-438-52-3913)

FEATURES  
source

1.6203  
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41.2255  
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/protein\_id="BAA20836.1"  
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LOHIGLNGQMAVAKRNETSAYIDQIGBLTFWFGSRGEKIKHAAATFCNSQPFAL  
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TEREVKKAADRCQIILFNQAVKAEKQKLEEDYORLDTSSLSLSSPVNEELRN  
LDLTRKMTIHGSLPVMKVRDKTIDLTYLLLEDIIVLLOKODRVLTRCHSKILASTA

DSKTFSEVILKSTVLVROVATDNKALFVISMDSNGAOIYELVAQTVSEKTVWQDILIC  
LIASVKEQSTPIPLPOSTPGEJNDDEDEPKLKEBOHGISVTGLQSPDRJLGEST  
RMSKPOSHSTSGSEVRLFEVABROPAKQHTDGLTKEVGEDYQJAIIPSHLPVS  
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IKAVHSGCHMFRTGTGDIATCYSPRTSTESFAPRDSVGLAPQDSQSNILYMHMTI  
MTPENPTHEPEGGLDSDGEHFPDAREAHSDENPSGSDGAVNKEEKDVNLIRISGNVLI  
DGYDVQSSSTDEEVASSLTLQPMGTGIPAVESTHQOQHSPOHTSHDGAISPTPEFLV  
OORWAMEVSCFEIQSPSSCADSQSQOIMEYIHKIEADLEHLKKEVESYTLICORLAGS  
ALTDKHSKDS"

BASE COUNT 1808 a 1195 c 1336 g 1864 t

ORIGIN

Query Match 80.0%; Score 16; DB 9; Length 6203;  
Best Local Similarity 100.0%; Pred. No. 1.5e+02;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 CTTAGAGGACACACT 17  
|||||  
Db 4168 CTTAGAGGACACACT 4183

RESULT 23

AF180681 9501 bp mRNA linear PRI 02-MAR-2000  
LOCUS Homo sapiens guanine nucleotide exchange factor (LARG) mRNA,  
DEFINITION complete cds.  
ACCESSION AF180681 GI:7110159  
VERSION AF180681.1 GI:7110159  
KEYWORDS Homo sapiens (human)  
SOURCE Homo sapiens  
ORGANISM Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.  
REFERENCE 1 (bases 1 to 9501)  
AUTHORS Koulas,P.J., Strout,M.P., Becknell,B., Veronese,M.L., Croce,C.M.,  
Thell,K.S., Krahe,R., Runtu,T., Knutti,L.S., Bloomfield,C.D. and  
Calligaris,M.A.  
TITLE Identification of a gene at 11q23 encoding a guanine nucleotide  
exchange factor: evidence for its fusion with MLL in acute myeloid  
leukemia  
JOURNAL Proc. Natl. Acad. Sci. U.S.A. 97 (5), 2145-2150 (2000)  
MEDLINE 20160919  
PUBMED 10681437  
REFERENCE 2 (bases 1 to 9501)  
AUTHORS Koulas,P.J., Strout,M.P., Becknell,B., Bloomfield,C.B. and  
Calligaris,M.A.  
TITLE Direct Submission  
JOURNAL Submitted (25-AUG-1999) Internal Medicine, Division of  
Hematology-Oncology, The Ohio State University, 4th floor  
Starling-Loving Hall, 320 West 10th Ave, Columbus, OH 43210, USA  
FEATURES  
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1. 9501  
Location/Qualifiers  
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/chromosome="11"  
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8. 4642  
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IKAVHSGCHMFRTGTGDIATCYSPRTSTESFAPRDSVGLAPQDSQSNILYMHMTI  
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ALTDKHSKDS"

VSQEDPATLLYISDLVKHTNSKETRIIFLEHFOFLDRSAHLKVSPDEMSADLEK  
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RLTEKEPCTEQIAKIEEVLMTQAVBEKSSYMOYVILMYMHGLKQVKEPRLND  
HKRGIRGLPYSKMKDKGEBEKGKRGKRPISIGPRRBRBNSAIGRAMLOKA  
RHRKRLSTPSSVSPERPOSAKLRQSGLANBEGDAGYLRANSMSSVAGSAPSEBQEGE  
NDTGSQVGESESAPEDTIDGTRTNTVDFPPPLDVOVECEVERTEHGTKRP  
RKPFSVAGSESQSEDEQENBLETDPNMQULVSEVLLGLPKCIKOEVLNELPYT  
ERAHVRLTKLVDOVFYQVRSREGILSPSELKIFSNLIDILQHLGLEKQVAKVKN  
ETSVIDOIGEDLTFWSPGEEKLKGHAATPCSNPALFEMIKSKQKDSRQFVOD  
AESNLICRLQDKDIIPTOMORLTYPILLNDIAKTYEMPTEREKVKAAADRCIOIN  
YVNOAVKEAKKORLEPYORRLDSLSLSEVPVNERLNDLTRKMHIEGELVWKV  
NNDKTIIDLTYLLEDETLVLQKODRLVFRKSKLSTABSKITFSEYIKLSTVLV  
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STPGEJNDDEDEPKLKEBOHGISVTGLQSPDRJLGESTLISKPOSHSTSGSE  
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LVQOLGLEKSVQEDMOPPRYPRTASQGPDDSVIOWSENIAKVSCHGHEMFPRTGSG  
DIATCYSPRTSTESFAPRDSVGLAPQDSQSNILYMHMTIIPREMPTEPEGGLDSDG  
EHFPAKHSNDENPSBGDAVNKEEKDVNLIRISGNVLIIDGYDVQSSSTDEEVASS  
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BASE COUNT 2816 a 1872 c 2078 g 2735 t

ORIGIN

Query Match 80.0%; Score 16; DB 9; Length 9501;  
Best Local Similarity 100.0%; Pred. No. 1.3e+02;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 CTTAGAGGACACACT 17  
|||||  
Db 6556 CTTAGAGGACACACT 6571

RESULT 24

AP000271 16529 bp DNA linear PRI 12-JUN-2003  
LOCUS Homo sapiens genomic DNA, chromosome 21 clone:CMF21-74A4, complete  
DEFINITION sequence.  
ACCESSION AP000271 GI:31621048  
VERSION AP000271.3 GI:31621048  
KEYWORDS HTG.  
SOURCE Homo sapiens (human)  
ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.  
REFERENCE 1  
AUTHORS Hattori,M., Ishii,K., Toyoda,A., Taylor,T.D., Hong-Seog,P.,  
Fujiyama,A., Yada,T., Totoki,Y., Watanabe,H. and Sakaki,Y.  
TITLE Homo sapiens genomic DNA  
JOURNAL Published Only in Database (1999)  
AUTHORS Hattori,M., Ishii,K., Toyoda,A., Taylor,T.D., Hong-Seog,P.,  
Fujiyama,A., Yada,T., Totoki,Y., Watanabe,H. and Sakaki,Y.  
TITLE Direct Submission  
JOURNAL Submitted (13-MAY-1999) Masahira Hattori, The Institute of Physical  
and Chemical Research (RIKEN), Genomic Sciences Center (GSC),  
1-7-22 Suenho-chou, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan  
(E-mail:hattori@gscc.riken.go.jp, URL:http://hgp.gsc.riken.go.jp/,  
Tel:81-45-503-9111, Fax:81-45-503-9170)  
COMMENT On Jun 12, 2003 this sequence version replaced gi:31071640.  
FEATURES  
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Db      5368 AGGAGACAAGTCCC 5383

RESULT 25
AF520762
LOCUS   AF520762
DEFINITION Homo sapiens X-ray repair complementing defective repair in Chinese hamster cells 2 (XRCC2) gene, complete cds.
ACCESSION AF520762
VERSION   AF520762.1 GI:21489901
KEYWORDS
SOURCE    Homo sapiens (human)
ORGANISM  Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
REFERENCE 1 (bases 1 to 32813)
AUTHORS  Rieder,M.J., Livingston,R.J., Braun,A.C., Montoya,M.A., Chung,M.-W., Miyamoto,K.E., Nguyen,C.P., Nguyen,D.A., Poel,C.L., Robertson,P.D., Schackwitz,W.S., Sherwood,J.K., Wittek,L.A. and Nickerson,D.A.
TITLE     Direct Submission
JOURNAL   Submitted (11-JUN-2002) Genome Sciences, University of Washington, 1705 NE Pacific, Seattle, WA 98195, USA
COMMENT   To cite this work please use: NIHES-SNPs, Environmental Genome Project, NIHES ES15478, Department of Genome Sciences, Seattle, WA (URL: http://egp.gs.washington.edu).
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964
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1133..1136
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1134
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2002
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Query Match 80.0%; Score 16; DB 9; Length 32813;  
Best Local Similarity 100.0%; Pred. No. 1e+02;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 5 AGAGAGACAAGTCCC 20  
DB 24451 AGAGAGACAAGTCCC 24466

RESULT 26

AC005203/c  
LOCUS AC005203 41308 bp DNA linear PRI 30-JUN-1998  
DEFINITION Homo sapiens chromosome 16, cosmid clone RT70 (LANL), complete  
sequence.  
ACCESSION AC005203  
VERSION AC005203.1 GI:3273381  
KEYWORDS HTG.  
SOURCE Homo sapiens (human)  
ORGANISM Homo sapiens  
REFERENCE Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.  
AUTHORS Kikkawa, D.O.  
TITLE Large Scale Sequence Analysis and Annotation with the Sequence  
Comparison Analysis (SCAN) System  
JOURNAL Unpublished  
REFERENCE 2 (bases 1 to 41308)  
AUTHORS Rikke, D.O., Bruce, D., Mundt, M., Doggett, N., Munk, C., Saunders, E.,  
Robinson, D., Jones, M., Buckingham, J., Chasteen, L., Thompson, S.,  
Goodwin, L., Bryant, J., Tesmer, J., Meincke, L., Longmire, J.,  
White, S., Ueng, S., Tatum, O., Campbell, C., Fawcett, J., Malbie, M.,  
Mistra, M. and Deaven, L.  
TITLE Sequencing of Human Chromosome 16p13.3  
JOURNAL Unpublished  
REFERENCE 3 (bases 1 to 41308)  
AUTHORS Rikke, D.O., Bruce, D., Mundt, M., Doggett, N., Munk, C., Saunders, E.,  
Robinson, D., Jones, M., Buckingham, J., Chasteen, L., Thompson, S.,  
Goodwin, L., Bryant, J., Tesmer, J., Meincke, L., Longmire, J.,  
White, S., Ueng, S., Tatum, O., Campbell, C., Fawcett, J., Malbie, M.,  
Mistra, M. and Deaven, L.  
TITLE Direct Submission  
JOURNAL Submitted (30-JUN-1998) Center for Human Genome Studies, DOE Joint  
Genome Institute, Los Alamos National Laboratory, MS M888, Los  
Alamos, NM 87545, USA  
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SOURCE Homo sapiens (human)  
 ORGANISM Homo sapiens  
 REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.  
 1 (bases 1 to 68589)  
 AUTHORS Birren, B., Linton, L., Nusbaum, C. and Lander, E.  
 TITLE Homo sapiens chromosome 18, clone RP11-886K22  
 JOURNAL Unpublished  
 REFERENCE 2 (bases 1 to 68589)  
 AUTHORS Birren, B., Linton, L., Nusbaum, C., Lander, E., Allen, N., Anderson, S., Barna, N., Baetien, V., Boguslavsky, L., Bouknight, B., Brown, A., Camarata, J., Campopiano, A., Choepel, Y., Colangelo, M., Collins, S., Collymore, A., Cooke, P., Dearellano, K., Dewar, K., Diaz, J. S., Dodge, S., Faro, S., Ferreira, P., Fitzhugh, W., Gage, D., Galagan, J., Gardy, S., Ginde, S., Goyette, M., Graham, L., Grand-pierre, N., Hagos, B., Heaford, A., Horton, L., Hulme, W., Iliev, I., Johnson, R., Jones, C., Karafas, A., Lacombe, K., Lamatares, R., Lander, T., Lehoczy, J., Levine, R., Liu, G., Maclean, C., MacDonald, P., Marquis, N., Mathews, C., McCarthy, M., McEwan, P., McKernan, K., McNetters, R., Meldrim, J., Meneus, L., Mihova, T., Mlenga, V., Murphy, T., Naylor, J., Nguyen, C., Nordu, C., Norman, C. H., O'Connor, T., O'Donnell, P., O'Neill, D., Oliver, T., Peterson, K., Phunhkhang, P., Pierre, N., Pollara, V., Raymond, C., Reta, R., Rieback, M., Riley, R., Rise, C., Rogov, P., Roman, J., Roselli, M., Roy, A., Santos, R., Schauer, S., Schupack, R., Seaman, S., Severy, P., Sougnez, C., Spencer, B., Stange-Thomann, N., Stojanovic, N., Straus, N., Subramanian, A., Talamas, J., Teafaye, S., Theodore, J., Travers, M., Travis, N., Triggilo, J., Vassiliev, H., Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Ye, W. J., Young, G., Zainoun, J., Zembek, L., Zimmer, A. and Zody, M.  
 TITLE Submitted (15-JAN-2001) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA  
 JOURNAL All repeats were identified using RepeatMasker:  
 COMMENT Smit, A.F.A. & Green, P. (1996-1997)  
 http://ftp.genome.washington.edu/RM/RepeatMasker.html  
 ----- Genome Center  
 Center: Whitehead Institute/ MIT Center for Genome Research  
 Web site: http://www-seq.wi.mit.edu  
 Contact: sequence\_submissions@genome.wi.mit.edu  
 ----- Project Information  
 Center project name: L12306  
 Center clone name: 886\_K\_22  
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 \* NOTE: This record contains 85 individual  
 \* sequencing reads that have not been assembled into  
 \* contigs. Runs of N are used to separate the reads  
 \* and the order in which they appear is completely  
 \* arbitrary. Low-pass sequence sampling is useful for  
 \* identifying clones that may be gene-rich and allows  
 \* overlap relationships among clones to be deduced.  
 \* However, it should not be assumed that this clone  
 \* will be sequenced to completion. In the event that  
 \* the record is updated, the accession number will  
 \* be preserved.  
 \* 1  
 \* 715 814: contig of 714 bp in length  
 \* 815 1522: contig of 708 bp in length  
 \* 1523 1622: gap of 100 bp  
 \* 1623 2330: contig of 708 bp in length  
 \* 2331 2430: gap of 100 bp  
 \* 2431 3132: contig of 702 bp in length  
 \* 3133 3232: gap of 100 bp  
 \* 3233 3944: contig of 712 bp in length  
 \* 3945 4044: gap of 100 bp  
 \* 4045 4741: contig of 697 bp in length  
 \* 4742 4841: gap of 100 bp  
 \* 4842 5570: contig of 729 bp in length  
 \* 5571 5670: gap of 100 bp  
 \* 5671 6387: contig of 717 bp in length  
 \* 6388 6487: gap of 100 bp  
 \* 6488 7189: contig of 702 bp in length

7190 7289: gap of 100 bp  
 7290 7992: contig of 703 bp in length  
 7993 8092: gap of 100 bp  
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 8897 9603: contig of 707 bp in length  
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 10506 11206: contig of 701 bp in length  
 11207 11306: gap of 100 bp  
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 12043 12142: gap of 100 bp  
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 14490 15269: gap of 100 bp  
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 16084 16183: gap of 100 bp  
 16184 16887: contig of 704 bp in length  
 16888 16987: gap of 100 bp  
 16988 17677: contig of 690 bp in length  
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 18584 19299: contig of 716 bp in length  
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 19400 20119: contig of 720 bp in length  
 20120 20219: gap of 100 bp  
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 20946 21045: gap of 100 bp  
 21046 21763: contig of 718 bp in length  
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 21864 22556: contig of 693 bp in length  
 22557 22656: gap of 100 bp  
 22657 23375: contig of 713 bp in length  
 23376 23475: gap of 100 bp  
 23476 24179: contig of 704 bp in length  
 24180 24279: gap of 100 bp  
 24279 24973: contig of 694 bp in length  
 24974 25073: gap of 100 bp  
 25074 25788: contig of 715 bp in length  
 25789 25888: gap of 100 bp  
 25889 26602: contig of 714 bp in length  
 26603 26702: gap of 100 bp  
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 27399 27498: gap of 100 bp  
 27499 28190: contig of 692 bp in length  
 28191 28290: gap of 100 bp  
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 29009 29108: gap of 100 bp  
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 29826 29925: gap of 100 bp  
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 30629 30728: gap of 100 bp  
 30729 31426: contig of 698 bp in length  
 31427 31526: gap of 100 bp  
 31527 32222: contig of 696 bp in length  
 32223 32322: gap of 100 bp  
 32323 33015: contig of 693 bp in length  
 33016 33115: gap of 100 bp  
 33116 33832: contig of 717 bp in length  
 33833 33932: gap of 100 bp  
 33933 34659: contig of 727 bp in length  
 34660 34759: gap of 100 bp  
 34760 35450: contig of 691 bp in length  
 35451 36270: gap of 100 bp  
 36270 36370: contig of 720 bp in length  
 36371 36370: gap of 100 bp

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* 36371 37078: contig of 708 bp in length
* 37079 37178: gap of 100 bp
* 37179 37892: contig of 714 bp in length
* 37893 37992: gap of 100 bp
* 37993 38715: contig of 723 bp in length
* 38716 38815: gap of 100 bp
* 38816 39528: contig of 713 bp in length
* 39529 39629: gap of 100 bp
* 39629 40340: contig of 712 bp in length
* 40341 40440: gap of 100 bp
* 40441 41084: contig of 644 bp in length
* 41085 41184: gap of 100 bp
* 41185 41877: contig of 693 bp in length
* 41878 41977: gap of 100 bp
* 41978 42655: contig of 678 bp in length
* 42656 42755: gap of 100 bp
* 42756 43564: contig of 709 bp in length
* 43565 44301: contig of 727 bp in length
* 44302 44401: gap of 100 bp
* 44402 45116: contig of 715 bp in length
* 45117 45216: gap of 100 bp
* 45217 45927: contig of 711 bp in length
* 45928 46027: gap of 100 bp
* 46028 46702: contig of 675 bp in length
* 46703 46802: gap of 100 bp
* 46803 47515: contig of 713 bp in length
* 47516 47615: gap of 100 bp
* 47616 48326: contig of 711 bp in length
* 48327 48426: gap of 100 bp
* 48427 49125: contig of 699 bp in length
* 49126 49225: gap of 100 bp
* 49226 49930: contig of 705 bp in length
* 49931 50030: gap of 100 bp
* 50031 50744: contig of 714 bp in length
* 50745 50844: gap of 100 bp
* 50845 51534: contig of 690 bp in length
* 51535 51634: gap of 100 bp
* 51635 52348: contig of 714 bp in length
* 52349 52448: gap of 100 bp
* 52449 53192: contig of 744 bp in length
* 53193 53292: gap of 100 bp
* 53293 53930: contig of 638 bp in length
* 53931 54030: gap of 100 bp
* 54031 54736: contig of 706 bp in length
* 54737 54836: gap of 100 bp
* 54837 55531: contig of 635 bp in length
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Query Match      80.0%; Score 16; DB 2; Length 68589;
Best Local Similarity 100.0%; Pred. No. 88;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Oy 5 AGGAGGAAACAGTCCC 20
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Db 63479 AGGAGGAAACAGTCCC 63464

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RESULT 28
AC003109 76850 bp DNA linear PRI 17-MAR-1998
LOCUS Human DNA from overlapping chromosome 7 PAC and PI clones
DEFINITION containing the XRCC2 gene, genomic sequence, complete sequence.
ACCESSION AC003109
VERSION AC003109.1 GI:2961444
KEYWORDS HTG.
SOURCE Homo sapiens (human)
ORGANISM Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 76850)
AUTHORS Liu,N., Lamerdin,J.E., Tebbes,R.S., Schild,D., Tucker,J.D., Shen,R.,
Brookman,K.W., Siciliano,M.J., Walter,C.A., Fan,W., Narayana,L.S.,

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/mol_type="genomic DNA"
/db_xref="taxon:9606"
/chromosome="7"
/map="7q36"
/clone="hXRCC2"
/note="PI clone 7515 obtained from Genome Systems (library
constructed from male foreskin-derived fibroblast line).
PAC clone 13620 obtained from Genome Systems (library
constructed from male leukocytes by P. de Jong)."
2. 208
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/complement(380..442)
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/complement(497..813)
/rpt_family="L1"
/complement(825..1106)
/rpt_family="Alu"
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2034..2157
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2731..3063
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4236..33903
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4322..31759
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SSBEIIKYCGRFLVYCSSTHLILTLVLSLEMPFSPHLCILIDLSLAFWIDY
NGESVYLOESTRKSCLEKLVNDRVLVPAITOTIMOKASSSSSEPSASRLCD
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VERC"
5618..5774
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repeat_region

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repeat_region complement (6948, .7208)
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repeat_region complement (8375, .8630)
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repeat_region complement (8887, .9129)
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repeat_region complement (9181, .9456)
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repeat_region complement (9515, .9795)
/rpt_family="Alu"
repeat_region complement (10534, .10574)
/rpt_family="Alu"
repeat_region complement (10709, .10968)
/rpt_family="Alu"
repeat_region complement (12008, .12076)
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repeat_region complement (12359, .12632)
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repeat_region complement (17529, .17829)
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repeat_region complement (18140, .18430)
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repeat_region complement (19259, .19503)
/rpt_family="Alu"
misc_feature /note="BLASTX similarity to (1088, .1146); match: 0.45,
score: 9.0e-06; database searched: nr; hypothetical
protein (L1H 3', region) - human"
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21931, .22256
/rpt_family="Alu"
repeat_region complement (22264, .22440)
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repeat_region complement (22820, .23070)
/rpt_family="Alu"
repeat_region complement (23153, .23436)
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repeat_region complement (24031, .24148)
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26621, .26930
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/rpt_family="Alu"
misc_feature complement (31037, .31372)
/rpt_family="Alu"
/gene="XRCC2"
/note="BLASTN similarity to Y08837 (1, .336); match: 0.99,
score: 7.0e-130; database searched: nr; H.sapiens mRNA for
RAD51-like protein-Other overlapping matches:
BLASTX similarity to PID (93, .149, 206, .243); match: 0.43,
score: 9.7e-06; database searched: nr; (Y13144) Rad51
homologue [Trapanosoma brucei]-BLASTX similarity to
2262211 (101, .162); match: 0.43, score: 1.8e-06; database
searched: nr; (U92068) RecA-like protein [Mus musculus]"
complement (31829, .32256)
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repeat_region complement (32533, .33613)
/rpt_family="L1"
32880, .33124
/note="predicted exon, program: gtail2exons_human_1.3,
frame: 1, quality: good, score: 67.000"
33635, .33888
/rpt_family="Alu"
repeat_region complement (33903, .34050)
/rpt_family="L1"
repeat_region complement (34480, .34609)
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repeat_region complement (35004, .35121)
/rpt_family="Alu"
misc_feature complement (36278, .36522)
/note="DBS similarity to AA255626 2831f06.s1 NCI-CGAP GCBI
Homo sapiens cDNA clone 666819 3' similar to contains Alu
repetitive element; contains element LTRB repetitive
element; (186, .427); 96% identity."
repeat_region complement (36396, .36646)
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repeat_region complement (36742, .37029)
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Query Match 80.0%; Score 16; DB 9; Length 76850;
Best Local Similarity 100.0%; Pred. No. 86;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 AGAGGAAACAAAGTCCC 20
Db 27146 AGAGGAAACAAAGTCCC 27161

RESULT 29
AX646415 84510 bp DNA linear PAT 04-MAR-2003
LOCUS Sequence 607 from Patent EP1270724.
DEFINITION AX646415
ACCESSION AX646415
VERSION AX646415.1 GI:28798796
KEYWORDS
SOURCE
ORGANISM Homo sapiens (human)
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE
AUTHORS Suwa,M., Asai,K., Akiyama,Y. and Aburatani,H.
TITLE Guanosine triphosphate-binding protein coupled receptors
JOURNAL Patent: EP 1270724-A 607 02-JAN-2003;
National Institute of Advanced Industrial Science and Technology
(JP) ; Center for Advanced Science and Technology Incubation, Ltd.

FEATURES
source Location/Qualifiers
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/organism="Homo sapiens"
/mol_type="genomic DNA"

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CDS
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      72348, .72417,75647, .75738,77542, .77668,82828, .82869,
      84170, .84310)
      /note="unnamed protein product"
      /codon_start=1
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      LFRNTOGWSETEPFPNLACGVNVNDSNRSHSYLKLVMYTVGSSSLWMLVA
      LGILCAFRLHCTRYRIIMHLFVSFIILEALNFIDAVLFSDPTVYCDARACKLV
      MLYRCCMANYSWLIVGLYLTLTASFESRKYLGFAPFGSGPAIYVALMAITRA
      RHPLEDSCMDINNASITWTIRGPVTLSIIINFPIINIRIKRIPOETRGENV
      SHVKRALSTLLIPFEITHIYVAFPSDEDMEIOLFELLAGSFQGLVAVLTCFLNA
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BASE COUNT      21253 a 19989 c 19504 g 23764 t

ORIGIN
Query Match      80.0%; Score 16; DB 6; Length 84510;
Best Local Similarity 100.0%; Pred. No. 84;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy      2 CTTAGGAGAACCAACT 17
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Db       66595 CTTAGGAGAACCAACT 66580

RESULT 30
AB065660/c      84510 bp DNA linear PRI 23-JUL-2002
LOCUS           Homo sapiens gene for seven transmembrane helix receptor, complete
DEFINITION      AB065660
VERSION         AB065660
KEYWORDS        Homo sapiens (human)
ORGANISM        Homo sapiens
AUTHORS         Mukaiyota, Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
                Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
TITLE           Suwa,M., Sato,T., Okouchi,I., Arita,M., Futami,K., Matsumoto,S.,
                Tezutsumi,S., Aburatani,H., Asai,K. and Akiyama,Y.
                Genome-wide discovery and analysis of human seven transmembrane
                helix receptor genes
JOURNAL         Unpublished
REFERENCE       2 (bases 1 to 84510)
AUTHORS         Suwa,M.
COMMENT         Direct Submission
                Submitted (11-JUL-2001) Makiko Suwa, Computational Biology Research
                Center (CBRC), National Institute of Advanced Industrial Science
                and Technology (AIST); 2-41-6 Aomii Koto-ku, Tokyo 135-0064, Japan
                (E-mail:m-suwa@aist.go.jp, URL:http://www.cbrc.jp/,
                Tel:81-3-3599-8080, Fax:81-3-3599-8081)
                This sequence is a seven transmembrane helix receptor candidate
                predicted from the whole human genome sequences using our automated
                system that contains programs of gene finding(GeneDecoder), sequence search, motif-domain assignment and
                transmembrane helix prediction.
                And the sequence is submitted by the collaborative project between
                [Computational Biology Research Center (CBRC), National Institute
                of Advanced Industrial Science and Technology (AIST)] and [Genome
                Science Division, Research Center for Advanced Science and
                Technology (RCAST), University of Tokyo].

FEATURES
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Best Local Similarity	100.0%	Pred. No. 84,							
Matches	16;	Conservative	0;	Mismatches	0;	Indels	0;	Gaps 0;	
Cy	2	CTTAGGAGAACCAACT	17						
Dd	66595	CTTAGGAGAACCAACT	66580						
RESULT 31									
LOCUS	ACOL10963	94357 bp	DNA	linear	HTG 01-OCT-1999				
DEFINITION	Homo sapiens clone 11_J_20, LOW-PASS SEQUENCE SAMPLING.								
ACCESSION	ACOL10963								
VERSION	ACOL10963.2	GI:606126							
KEYWORDS	HTG; HTGS PHASED.								
SOURCE	Homo sapiens (human)								
ORGANISM	Homo sapiens								
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;								
AUTHORS	Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.								
TITLE	1 (bases 1 to 94357)								
JOURNAL	Biren,B., Linton,L., Nusbaum,C. and Lander,E.								
REFERENCE	Homo sapiens, clone 11_J_20								
AUTHORS	Unpublished								
	2 (bases 1 to 94357)								
	Biren,B., Linton,L., Nusbaum,C., Lander,E., Allen,N., Anderson,M.,								
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	Brown,A., Castle,J.A., Colangelo,M., Collins,S., Collimore,A.,								
	Cooke,P., Daretellano,K., Dewar,K., Domino,M., Donelan,L., Doyle,M.,								
	Ferreira,P., Fitzhugh,W., Forrest,C., Funke,R., Gage,D.,								
	Galagan,J., Gardyna,S., Grant,G., Hagos,B., Heatford,A., Horton,L.,								
	Howland,J.C., Johnson,R., Jones,C., Kann,L., Karatas,A., Klein,J.,								
	Lehoucq,J., Lieu,C., Locke,K., Macdonald,P., Margush,N.,								
	Melwan,P., McGurk,A., McKernan,K., McLaughlin,J., Meldrum,J.,								
	Morrow,J., Naylor,J., Norman,C.H., O'Connor,T., O'Donnell,P.,								
	Peterson,K., Pollara,V., Riley,R., Roy,A., Santos,R., Severy,P.,								
	Stange-Thomann,N., Stojanovic,N., Subramanian,A., Talamas,J.,								
	Teifayez,S., Titrell,A., Vassiliev,H., Vo,A., Wheeler,J., Wu,X.,								
	Wyman,D., Ye,W.J., Zimmer,A. and Zody,M.								
TITLE	Direct Submission								
JOURNAL	Submitted (28-SEP-1999) Whitehead Institute/MIT Center for Genome								
COMMENT	Research, 320 Charles Street, Cambridge, MA 02141, USA								
	On Oct 1, 1999 this sequence version replaced gi:5931399.								
	All repeats were identified using RepeatMasker: Smit,A.F.A. &								
	Green,P. (1996-1997)								
	http://fhp.genome.washington.edu/RM/RepeatMasker.html.								
	* NOTE: This record contains 98 individual								
	* sequencing reads that have not been assembled into								
	* contigs. Runs of N are used to separate the reads								
	* and the order in which they appear is completely								
	* arbitrary. Low-pass sequence sampling is useful for								
	* identifying clones that may be gene-rich and allows								
	* overlap relationships among clones to be deduced.								
	* However, it should not be assumed that this clone								

\* will be sequenced to completion. In the event that  
\* the record is updated, the accession number will  
\* be preserved.  
1 892: contig of 892 bp in length  
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Query Match	Query Local Similarity	Score	DB	Length	100000;
Matches 16; Conservative	100.0%; Pred. No. 81; Mismatches 0; Indels 0; Gaps 0;	80.0%; Score 16; DB 9; Length 100000;			

Db	11338	ACGAGGAAACAAGTCCC	113323
RESULT 33	AP000104	100000 bp	DNA linear PRI 25-MAY-2002
LOCUS	AP000104		
DEFINITION	Homo sapiens genomic DNA of 21q22.1, GART and AML related,		
ACCESSION	OT8C10-149C3	region, segment 7/20.	
VERSION	AP000104		
KEYWORDS	AP000104.1	GI:4730838	
SOURCE	Homo sapiens (human)		
ORGANISM	Homo sapiens		
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;		
AUTHORS	Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.		
TITLE	Hattori,M., Ishii,K., Toyoda,A., Shiba,T. and Sakaki,Y.		
JOURNAL	Homo sapiens 2,051,516bp genomic DNA of 21q22.1 GART and AML region		
REFERENCE	Published Only in Database (1999)		
AUTHORS	2 (bases 1 to 100000)		
TITLE	Hirakawa,M., Yamaguchi,H., Imai,K. and Shimada,J.		
JOURNAL	Direct Submission		
	Submitted (15-APR-1999) Mika Hirakawa, Japan Science and Technology		
	Corporation (JST), Advanced Databases Department, 5-3, Yonbancho,		
	Chiyoda-ku, Tokyo 102-0081, Japan (E-mail:mika@ccyo.jst.go.jp,		
	URL:http://www.alls.tokyo.jst.go.jp/, Tel:81-3-5214-8491,		
	Fax:81-3-5214-8470)		
COMMENT	This sequence is conducted by Kitasato University JST sequencing		
	Laboratory as a JST sequencing team.		
	Principal Investigator:Yoshiyuki Sakaki Ph.D.		
	Phone:+81-3-5449-5622, Fax : +81-3-5449-5445,		
	sakaki@cc.yms.u-tokyo.ac.jp		
	Sub-leader: Tadayoshi Shiba Ph.D., Maehira Hattori Ph.D. The		
	sequence is submitted by Human Genome Sequencing in ALIS project of		
	JST.		
	Japan Science and Technology Corporation (JST)		
	5-3, Yonbancho, Chiyoda-ku, Tokyo 102-0081 Japan		
	For further information about this sequence, including its location		
	and relationship to other sequences, please visit our sequence		
	archive Web site (http://www.alls.tokyo.jst.go.jp/HGS/) or send		
	email to webmaster@www.alls.tokyo.jst.go.jp.		
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STS	/standard_name="D21S1413"		
	/note="UT7582;The location is between each flanking site		
	of PCR primers."		
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STS	/note="SHGC-51923;The location is between each flanking		
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BASE COUNT
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Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY
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    |||||
    25561 AGGAGGACAAGTCCC 25576

RESULT 34
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LOCUS
DEFINITION
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    clone 078C10-f32E9, segment 7/21, complete sequence.
ACCESSION
    AF000180
VERSION
    AF000180.1 GI:4827079
KEYWORDS
    HTG.
SOURCE
    Homo sapiens (human)
ORGANISM
    Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
    Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE
    1 (bases 1 to 100000)
    Hattori,M., Ishii,K., Toyoda,A., Taylor,T.D., Hong-Seog,P.,
    Fujiyama,A., Yada,T., Totoki,Y. and Sakaki,Y.
    Homo sapiens 2,083,744bp genomic DNA of 21q22.1 (REGION:
    D21S226-AML CLONE RANGER: Q78C10-f32E9)
    Published Only in Database (1999)
    2 (bases 1 to 100000)
    Hattori,M., Ishii,K., Toyoda,A., Taylor,T.D., Hong-Seog,P.,
    Fujiyama,A., Yada,T., Totoki,Y. and Sakaki,Y.
    Direct Submission
    Submitted (10-MAY-1999) Masahira Hattori, The Institute of Physical
    and Chemical Research (RIKEN), Genomic Sciences Center (GSC),
    Kitasato Univ., 1-15-1 Kitasato, Sagamihara, Kanagawa 228-8555,
    Japan (E-mail:hattori@gsc.riken.go.jp,
    URL:http://hgp.gsc.riken.go.jp/, Tel:81-42-778-9923,
    Fax:81-42-778-9924)

FEATURES
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BASE COUNT
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ORIGIN

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Query Match
Best Local Similarity 100.0%; Score 16; DB 9; Length 100000;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY
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    |||||
    25561 AGGAGGACAAGTCCC 25576

RESULT 35
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LOCUS
DEFINITION
    Homo sapiens chromosome 11 clone CMB9-2L13 map 11q23, WORKING DRAFT
    SEQUENCE, 13 unordered pieces.
ACCESSION
    AP000681.3 GI:8118869
VERSION
    HTG; HTGS_PHASE1; HTGS_DRAFT.
KEYWORDS
    HTG; HTGS_PHASE1; HTGS_DRAFT.
SOURCE
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ORGANISM
    Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
    Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE
    1 (bases 1 to 109149)
    Hattori,M., Ishii,K., Toyoda,A., Taylor,T.D., Hong-Seog,P.,
    Fujiyama,A., Yada,T., Totoki,Y., Watanabe,H. and Sakaki,Y.
    Homo sapiens 109,149 genomic DNA of 11q23
    Published Only in Database (1999)
    2 (bases 1 to 109149)
    Hattori,M., Ishii,K., Toyoda,A., Taylor,T.D., Hong-Seog,P.,
    Fujiyama,A., Yada,T., Totoki,Y., Watanabe,H. and Sakaki,Y.
    Direct Submission
    Submitted (08-NOV-1999) Masahira Hattori, The Institute of Physical
    and Chemical Research (RIKEN), Genomic Sciences Center (GSC),
    Kitasato Univ., 1-15-1 Kitasato, Sagamihara, Kanagawa 228-8555,
    Japan (E-mail:hattori@gsc.riken.go.jp,
    URL:http://hgp.gsc.riken.go.jp/, Tel:81-42-778-9923,
    Fax:81-42-778-9924)
    On May 31, 2000 this sequence version replaced gi:6997555.

COMMENT
    ----- Genome Center
    Center: RIKEN Genomic Sciences Center(GSC)
    Center code: RIKEN
    Web site: http://hgp.gsc.riken.go.jp/
    Contact: hattori@gsc.riken.go.jp
    ----- Project Information
    Center project name: Humberd11
    Center clone name: CMB9-2L13
    ----- Summary Statistics
    Sequencing vector: PCR products; 100% of reads
    Chemistry: Dye-terminator ET-amersham; 100% of reads
    Assembly program: Phrap; version 0.990329
    Consensus quality: 99833 bases at least Q40
    Consensus quality: 104437 bases at least Q30
    Consensus quality: 106696 bases at least Q20
    Insert size: 107949; sum-of-contigs
    Quality coverage: 4.90x in Q20 bases; sum-of-contigs
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    NOTE: This is a 'working draft' sequence. It currently consists of
    13 contigs. The true order of the pieces is not known and their
    order in this sequence record is arbitrary. Gaps between the
    contigs are represented as runs N, but the exact sizes of the gaps
    are unknown. This record will be updated with the finished sequence
    as soon as it is available and the accession number will be
    preserved
    1
    15418 contig of 15418 bp in length
    15519
    32589 contig of 17071 bp in length
    32690
    46533 contig of 13844 bp in length
    46634
    59641 contig of 13068 bp in length
    59742
    69969 contig of 10228 bp in length
    70070
    79400 contig of 9331 bp in length
    79501
    86689 contig of 7189 bp in length
    86790
    94162 contig of 7373 bp in length
    94263
    100141 contig of 5879 bp in length
    100242
    104375 contig of 4134 bp in length

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104476 107052 contig of 2577 bp in length  
 108671 contig of 1519 bp in length  
 107153 109149 contig of 378 bp in length  
 108772  
 Sequence updated (01-Feb-2000)  
 Sequence updated (26-May-2000).  
 \* NOTE: This is a 'working draft' sequence. It currently  
 \* consists of 13 contigs. The true order of the pieces  
 \* is not known and their order in this sequence record is  
 \* arbitrary. Gaps between the contigs are represented as  
 \* runs of N, but the exact sizes of the gaps are unknown.  
 \* This record will be updated with the finished sequence  
 \* as soon as it is available and the accession number will  
 \* be preserved.

1 15418: contig of 15418 bp in length  
 \* 15419 15518: gap of 100 bp  
 \* 15519 32589: contig of 17071 bp in length  
 \* 32590 32689: gap of 100 bp  
 \* 32690 46533: contig of 13844 bp in length  
 \* 46534 46633: gap of 100 bp  
 \* 46634 59641: contig of 13008 bp in length  
 \* 59642 59741: gap of 100 bp  
 \* 59742 69969: contig of 10228 bp in length  
 \* 69970 70069: gap of 100 bp  
 \* 70070 79400: contig of 9331 bp in length  
 \* 79401 79500: gap of 100 bp  
 \* 79501 86689: contig of 7189 bp in length  
 \* 86690 86790: gap of 100 bp  
 \* 86790 94262: contig of 7373 bp in length  
 \* 94263 94263: gap of 100 bp  
 \* 94264 100141: contig of 5879 bp in length  
 \* 100142 100241: gap of 100 bp  
 \* 100242 104375: contig of 4134 bp in length  
 \* 104376 104476: gap of 100 bp  
 \* 104476 107052: contig of 2577 bp in length  
 \* 107053 107153: gap of 100 bp  
 \* 107153 108671: contig of 1519 bp in length  
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 Location/Qualifiers  
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## FEATURES

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## ORIGIN

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 Best Local Similarity 100.0%; Pred. No. 79;  
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 CTTAGGAGAACAACT 17

DB 73711 CTTAGGAGAACAACT 73726

Search completed: August 15, 2003, 09:33:56  
 Job time : 555.75 secs

GenCore version 5.1.6  
Copyright (c) 1993 - 2003 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: August 14, 2003, 21:41:37 ; Search time 1252 Seconds

(without alignments)  
388.250 Million cell updates/sec

Title: US-10-074-620-2

Perfect score: 20

Sequence: 1 cctaagaggaacacagcacc 20

Scoring table: OLIGO\_NUC

Gapop 60.0 , Gapext 60.0

Searched: 22781392 seqs, 12152238056 residues

Word size : 0

Total number of hits satisfying chosen parameters: 45562784

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Listing first 120 summaries

Database :

EST:\*  
1: em\_estdb:\*  
2: em\_esthm:\*  
3: em\_estin:\*  
4: em\_estmu:\*  
5: em\_estcov:\*  
6: em\_estpl:\*  
7: em\_estcro:\*  
8: em\_hic:\*  
9: gb\_est1:\*  
10: gb\_est2:\*  
11: gb\_hic:\*  
12: gb\_est3:\*  
13: gb\_est4:\*  
14: gb\_est5:\*  
15: em\_estfun:\*  
16: em\_estom:\*  
17: em\_gss\_hum:\*  
18: em\_gss\_inv:\*  
19: em\_gss\_pln:\*  
20: em\_gss\_vit:\*  
21: em\_gss\_fun:\*  
22: em\_gss\_mam:\*  
23: em\_gss\_mus:\*  
24: em\_gss\_pro:\*  
25: em\_gss\_rtd:\*  
26: em\_gss\_phg:\*  
27: em\_gss\_vit:\*  
28: gb\_gss1:\*  
29: gb\_gss2:\*

Pred. No. is the number of results predicted by chance to have a  
score greater than or equal to the score of the result being printed,  
and is derived by analysis of the total score distribution.

# SUMMARIES

Result	Score	Query	Match	Length	DB	ID	Description
1	17	85.0	676	10	BE964731	601658274	
2	17	85.0	817	10	BF142291	601791832	
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4	16	80.0	203	10	BE755758	BE755758 209682 MA	

5	16	80.0	242	14	Z42603		Z42603 HSCOPA091 n
6	16	80.0	350	9	AU231688		AU231688 AU231688
7	16	80.0	358	14	R24852		R24852 YG29E05.r1
8	16	80.0	390	28	AQ483201		AQ483201 RPCT-11-2
9	16	80.0	395	5	A1696726		A1696726 wC56d11.x
10	16	80.0	400	14	T75305		T75305 YC89C08.r1
11	16	80.0	413	28	AQ379270		AQ379270 RPCT11-15
12	16	80.0	431	9	AM769407		AM769407 h16g12.x
13	16	80.0	443	10	BE273333		BE273333 894030E10
14	16	80.0	464	13	BY414889		BY414889 BY414889
15	16	80.0	476	14	R24326		R24326 YG32E03.r1
16	16	80.0	487	10	BF830043		BF830043 MR3-HN005
17	16	80.0	494	14	R24340		R24340 YG32H03.r1
18	16	80.0	516	10	BB693705		BB693705 BB693705
19	16	80.0	516	12	BG994855		BG994855 PMO-HT116
20	16	80.0	524	14	CA405555		CA405555 1001704.H
21	16	80.0	550	13	BQ347567		BQ347567 CM0-HT017
22	16	80.0	554	14	W52429		W52429 zc94E08.r1
23	16	80.0	591	12	BI751017		BI751017 T401_05d0
24	16	80.0	599	2	HS0095392		BK501675 Homo sapi
25	16	80.0	615	9	AU137185		AU137185 AU137185
26	16	80.0	620	12	BI156614		BI156614 602921206
27	16	80.0	661	9	AV821593		AV821593 AV821593
28	16	80.0	668	12	BU273261		BU273261 BU273261
29	16	80.0	703	13	BQ539794		BQ539794 PTAM0102
30	16	80.0	740	13	BU635192		BU635192 003D06.In
31	16	80.0	745	12	BI115344		BI115344 602863159
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33	16	80.0	872	12	BI088175		BI088175 602851213
34	16	80.0	879	13	BU177008		BU177008 AGENCOURT
35	16	80.0	884	10	BG740602		BG740602 602631028
36	16	80.0	886	10	BG501889		BG501889 602548991
37	16	80.0	900	13	BU509687		BU509687 AGENCOURT
38	16	80.0	904	10	BE869754		BE869754 601445669
39	16	80.0	942	13	BQ225365		BQ225365 AGENCOURT
40	16	80.0	966	13	BQ277316		BQ277316 AGENCOURT
41	16	80.0	978	9	AL536291		AL536291 AL536291
42	16	80.0	1201	9	AL546318		AL546318 AL546318
43	16	75.0	179	13	BY607541		BY607541 BY607541
44	15	75.0	193	28	BZ181647		BZ181647 CH230-340
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46	15	75.0	280	10	BE055057		BE055057 GA_Ea003
47	15	75.0	286	10	BB450936		BB450936 BB450936
48	15	75.0	287	10	BE054889		BE054889 GA_Ea002
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51	15	75.0	307	28	BH071600		BH071600 RPCT-24-2
52	15	75.0	315	10	BB317883		BB317883 BB317883
53	15	75.0	372	9	AA155221		AA155221 mr97C03.r
54	15	75.0	372	9	AA155224		AA155224 mr97C03.r
55	15	75.0	380	9	AA155220		AA155220 mr97C03.r
56	15	75.0	387	13	BY155062		BY155062 BY155062
57	15	75.0	413	9	AA155222		AA155222 mr97C04.r
58	15	75.0	421	28	AZ318410		AZ318410 1M0037P14
59	15	75.0	437	9	AA546043		AA546043 vk61C08.r
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63	15	75.0	460	14	N80106		N80106 YZ87F02.r1
64	15	75.0	460	14	N80119		N80119 YZ87F02.r1
65	15	75.0	460	28	AZ802618		AZ802618 2M0061003
66	15	75.0	475	28	AZ943693		AZ943693 2M0204022
67	15	75.0	477	10	BE270306		BE270306 GA_EB000
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73	15	75.0	499	29	BZ925689		BZ925689 CH240_72D
74	15	75.0	505	28	AZ597111		AZ597111 1M0410113
75	15	75.0	510	9	AJ225437		AJ225437 AJ225437
76	15	75.0	519	9	AV856135		AV856135 AV856135
77	15	75.0	523	28	AZ015440		AZ015440 RPCT-23-3

78	15	75.0	524	10	BC440460	GA_Ea000
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C 80	15	75.0	538	28	A0558274	HS 2066 B
C 81	15	75.0	542	14	CB873512	K0325G05
C 82	15	75.0	555	13	BQ405699	GA_Ed008
C 83	15	75.0	555	13	BQ414368	GA_Ed008
C 84	15	75.0	556	28	A2079907	RPCT-23-4
C 85	15	75.0	560	28	A2625916	1M0465C24
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C 87	15	75.0	566	28	A2792636	2M0045C15
C 88	15	75.0	573	28	A2625800	1M0465F12
C 89	15	75.0	575	9	AV880832	AV880832
C 90	15	75.0	577	13	B0694428	LL21n1197
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C 92	15	75.0	592	28	A0462877	HS 5212 A
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C 94	15	75.0	597	13	B0608042	BRY_3943
C 95	15	75.0	601	9	AA422164	2V31C01.r
C 96	15	75.0	606	10	BE288544	601094160
C 97	15	75.0	607	29	AG150407	Pan tce91
C 98	15	75.0	609	9	AI927904	wp03c07.x
C 99	15	75.0	611	29	BX133940	Danio rer
C 100	15	75.0	617	10	BG441695	GA_Ea001
C 101	15	75.0	617	28	A261932	RPCT-23-1
C 102	15	75.0	620	10	BG442675	GA_Ea001
C 103	15	75.0	626	9	AV880047	AV880047
C 104	15	75.0	626	28	BH059230	RPCT-24-3
C 105	15	75.0	628	13	BQ410892	GA_Ed003
C 106	15	75.0	630	13	BQ413572	GA_Ed007
C 107	15	75.0	634	10	BE985077	UT-M-CGDP
C 108	15	75.0	636	13	BQ410893	GA_Ed001
C 109	15	75.0	636	10	BG442578	GA_Ea001
C 110	15	75.0	640	13	BQ407961	GA_Ed000
C 111	15	75.0	640	13	BQ414737	GA_Ed009
C 112	15	75.0	641	28	A2113409	RPCT-23-4
C 113	15	75.0	643	9	A1585535	VK61c08.Y
C 114	15	75.0	645	9	AV889725	AV889725
C 115	15	75.0	646	9	AV856437	AV856437
C 116	15	75.0	646	28	A2255189	RPCT-23-1
C 117	15	75.0	650	28	A2458615	1M0262E23
C 118	15	75.0	651	13	BQ413525	GA_Ed007
C 119	15	75.0	652	10	BG442693	GA_Ea001
C 120	15	75.0	653	10	BG072204	H3107H11-

## ALIGNMENTS

RESULT 1  
BE964731 676 bp mRNA linear EST 14-DEC-2000  
LOCUS 601658274R1 NIH\_MGC\_69 Homo sapiens cDNA clone IMAGE:3885642 3',  
DEFINITION mRNA sequence.  
ACCESSION BE964731  
VERSION BE964731.2 GI:11768351  
KEYWORDS EST.  
SOURCE Homo sapiens (human)  
ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.  
REFERENCE 1 (bases 1 to 676)  
NIH-MGC http://mhc.nci.nih.gov/  
National Institutes of Health, Mammalian Gene Collection (MGC)  
Unpublished  
On Oct 3, 2000 this sequence version replaced gi:10575436.  
Contact: Robert Strausberg, Ph.D.  
Email: cgaabbs-remail.nih.gov  
Tissue Procurement: DCTD/DTF/Gazdar  
cDNA Library Preparation: Life Technologies, Inc.  
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNLN)  
DNA Sequencing by: Incyte Genomics, Inc.  
Clone distribution: MGC clone distribution information can be  
found through the I.M.A.G.E. Consortium/LNLN at:

http://image.lnl.gov  
Plate: L1CM648 row: c column: 19  
High quality sequence stop: 227.  
Location/Qualifiers  
1..676  
/organism="Homo sapiens"  
/mol\_type="mRNA"  
/db\_xref="taxon:9606"  
/clone="IMAGE:3885642"  
/tissue\_type="large cell carcinoma, undifferentiated"  
/lab\_host="DH10B (phage-resistant)"  
/clone\_lib="NIH\_MGC\_69"  
/note="Organ: Lung; Vector: pCMV-SPORT6; Site\_1: NotI;  
Site\_2: SalI; Cloned unidirectionally. Primer: Oligo dt.  
Average insert size 1.1 kb. Library constructed by Life  
Technologies."

BASE COUNT 139 a 138 c 193 g 206 t  
ORIGIN

Query Match 85.0%; Score 17; DB 10; Length 676;  
Best Local Similarity 100.0%; Pred. No. 23;  
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 TTAGGAGGACACTCC 19  
|||||  
Db 436 TTAGGAGGACACTCC 452

## RESULT 2

BF142291/c 817 bp mRNA linear EST 24-OCT-2000  
LOCUS 601791832P1 NCI\_CGAP\_Lu30 Mus musculus cDNA clone IMAGE:4022647 5',  
DEFINITION mRNA sequence.  
ACCESSION BF142291  
VERSION BF142291.1 GI:10981241  
KEYWORDS EST.  
SOURCE Mus musculus (house mouse)  
ORGANISM Mus musculus  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.  
REFERENCE 1 (bases 1 to 817)  
NIH-MGC http://mhc.nci.nih.gov/  
National Institutes of Health, Mammalian Gene Collection (MGC)  
Unpublished  
Contact: Robert Strausberg, Ph.D.  
Email: cgaabbs-remail.nih.gov  
Tissue Procurement: Gilbert Smith, Ph.D.  
cDNA Library Preparation: Life Technologies, Inc.  
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNLN)  
DNA Sequencing by: Incyte Genomics, Inc.  
Clone distribution: NCI-CGAP clone distribution information can be  
found through the I.M.A.G.E. Consortium/LNLN at:  
http://image.lnl.gov  
Plate: L1AM9279 row: p column: 08  
High quality sequence stop: 672.  
Location/Qualifiers  
1..817  
/organism="Mus musculus"  
/mol\_type="mRNA"  
/strain="C57BL/6J"  
/db\_xref="taxon:10090"  
/clone="IMAGE:4022647"  
/tissue\_type="tumor, metastatic to mammary"  
/lab\_host="DH10B"  
/clone\_lib="NCI\_CGAP\_Lu30"  
/note="Organ: Lung; Vector: pCMV-SPORT6; Site\_1: NotI;  
Site\_2: SalI; transgenic model WNT-1, expression driven by  
MMTV-LTR enhancer; Cloned unidirectionally. Primer: Oligo  
dt. Library constructed by Life Technologies.  
Investigator providing samples: Gilbert Smith, NIH"

## FEATURES

source

BASE COUNT 186 a 237 c 201 g 193 t  
ORIGIN



Query Match 85.0%; Score 17; DB 10; Length 817;  
 Best Local Similarity 100.0%; Pred. No. 24;  
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CCTAGAGGAGAACAGT 17  
 |||||  
 Db 773 CCTAGAGGAGAACAGT 757

RESULT 3  
 BM751719 200 bp mRNA linear EST 04-MAR-2002  
 LOCUS BM751719  
 DEFINITION K-EST0027999 S9SNU601 Homo sapiens CDNA clone S9SNU601-24-D05 5',  
 mRNA sequence.

ACCESSION BM751719 GI:19081351  
 VERSION  
 KEYWORDS  
 SOURCE  
 ORGANISM

Homo sapiens (human)  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.

REFERENCE 1 (bases 1 to 200)  
 Kim,N.S., Hahn,Y., Oh,J.H., Lee,J.Y., Ahn,H.Y., Chu,M.Y., Kim,M.R.,  
 Oh,K.U., Cheong,J.E., Sohn,H.Y., Kim,J.M., Park,H.S., Kim,S. and  
 Kim,Y.S.

TITLE 21C Frontier Korean EST Project 2001  
 JOURNAL  
 COMMENT

Genome Research Center  
 Korea Research Institute of Bioscience & Biotechnology  
 52 Boeun-dong Yuseong-gu, Daejeon 305-333, South Korea  
 Tel: +82-42-860-4470  
 Fax: +82-42-860-4409  
 Email: yongsung@mail.kribb.re.kr  
 Plate: 24 row: D column: 05  
 High quality sequence stop: 200.

FEATURES  
 source 1..200  
 Location/Qualifiers

1..200  
 /organism="Homo sapiens"  
 /mol\_type="mRNA"  
 /db\_xref="taxon:9606"  
 /clone="S9SNU601-24-D05"  
 /sex="M"  
 /tissue\_type="Ascites"  
 /cell\_type="Epithelial"  
 /lab\_host="SNU-601"  
 /lab\_host="Top10F"  
 /clone\_id="S9SNU601"  
 /note="Organ: Stomach; Vector: pME18-FL3; Site 1: XhoI;  
 Site 2: XhoI; The poly (A) + RNA was dephosphorylated with  
 bacterial alkaline phosphatase (BAP) and then dephosphorylated  
 with tobacco acid pyrophosphatase (TAP). The dephosphorylated  
 intact mRNA was ligated with DNA-RNA linker including SfiI  
 site by treatment of T4 RNA ligase and the first strand  
 cDNA was synthesized with Superscript II using SfiI  
 oligo-dT primer. After first strand synthesis, RNA was  
 degraded by NaOH treatment and cDNA was amplified by PCR  
 reaction. The PCR products were digested with SfiI and  
 cloned into DraIII-digested pME18-FL3 vector. The  
 obtained cDNA vectors were used for transformation of  
 competent cells E. coli Top10F by electroporation method.  
 The cDNA libraries constructed by this method are  
 full-length enriched cDNA library."

BASE COUNT 73 a 34 c 37 g 56 t  
 ORIGIN

Query Match 80.0%; Score 16; DB 12; Length 200;  
 Best Local Similarity 100.0%; Pred. No. 65;  
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 2 CTTAGAGGAGAACAGT 17  
 |||||  
 Db 92 CTTAGAGGAGAACAGT 107

RESULT 4  
 BE755758 203 bp mRNA linear EST 25-APR-2001  
 LOCUS BE755758  
 DEFINITION 209682 MARC 2BOV Bos taurus CDNA 5', mRNA sequence.

ACCESSION BE755758  
 VERSION BE755758.1 GI:10169750  
 KEYWORDS  
 SOURCE  
 ORGANISM

Bos taurus (cow)  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;  
 Bovidae; Bos.

REFERENCE 1 (bases 1 to 203)  
 Smith,T.P.L., Grose,W.M., Freking,B.A., Roberts,A.J., Stone,R.T.,  
 Casas,E., Wray,J.E., White,J., Cho,J., Fahrenkrug,S.C., Bennett  
 G.L., Heaton,M.P., Laegreid,W.W., Rohrer,G.A., Chitko-Mckown,C.G.,  
 Pertea,G., Holt,I., Karamycheva,S., Liang,F., Quackenbush,J. and  
 Keefe,J.W.

Sequence evaluation of four pooled-tissue normalized bovine cDNA  
 libraries and construction of a gene index for cattle  
 Genome Res. 11 (4), 626-630 (2001)

TITLE  
 JOURNAL  
 MEDLINE  
 PUBMED  
 COMMENT

Contact: Smith TPL  
 USDA, ARS, US Meat Animal Research Center  
 PO Box 166, Clay Center, NE 68933-0166, USA  
 Tel: 402 762 4366  
 Fax: 402 762 4390

Email: smitht@mail.marc.usda.gov  
 Single pass sequencing. Bases called and alt trimmed with phred  
 v0.980904.e. Vector identified by cross\_match with the -minscore 18  
 and -mismatch 12 options.

PCR primers  
 FORWARD: AGGAACAGCTATGACCAT  
 BACKWARD: GTTTCACGTCACGACG  
 Plate: 59 row: L column: 7  
 Seq primer: ATTTAGGAGACATATAG.

FEATURES  
 source 1..203  
 Location/Qualifiers

1..203  
 /organism="Bos taurus"  
 /mol\_type="mRNA"  
 /db\_xref="taxon:9913"  
 /tissue\_type="pooled"  
 /lab\_host="DH10B"  
 /clone\_id="MARC 2BOV"  
 /note="Vector: pCMV SPORT6; Site 1: NotI; Site 2: SalI;  
 library made from pooled tissue from testis, thymus,  
 semitendinosus muscle, longissimus muscle, pancreas,  
 adrenal, and endometrium."

BASE COUNT 70 a 29 c 48 g 56 t  
 ORIGIN

Query Match 80.0%; Score 16; DB 10; Length 203;  
 Best Local Similarity 100.0%; Pred. No. 65;  
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 2 CTTAGAGGAGAACAGT 17  
 |||||  
 Db 102 CTTAGAGGAGAACAGT 117

RESULT 5  
 Z42603 242 bp mRNA linear EST 10-NOV-1994  
 LOCUS Z42603  
 DEFINITION HSCOPA091 normalized infant brain CDNA Homo sapiens CDNA clone

c-09a09, mRNA sequence.  
 ACCESSION Z42603 GI:567353  
 VERSION  
 KEYWORDS  
 SOURCE  
 ORGANISM

Homo sapiens (human)  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.

REFERENCE  
AUTHORS  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.  
1 (bases 1 to 242)

TITLE  
JOURNAL  
MEDLINE  
PUBMED  
95277534  
7757816  
Contact: Genethon  
Genexpress-Genethon  
Genethon Centre de recherche sur le Genome Humain  
1, rue de l'Internationale, BP60 91002 EVRY Cedex, FRANCE  
Tel: 33169472800  
Fax: 33160778698  
Email: genexpress@genethon.fr  
Single read.  
Genexpress library id: C; Genexpress\_sequence\_id: y1c-0pa09  
Seq primer: (-21)M13 universal.  
Location/Qualifiers

FEATURES  
source  
1..242  
/organism="Homo sapiens"  
/mol\_type="mRNA"  
/db\_xref="taxon:9606"  
/clone="c-0pa09"  
/sex="Female"  
/tissue\_type="total brain"  
/dev\_stage="3 months old"  
/clone\_lib="normalized infant brain cDNA"  
/note="Organ: brain; Vector: lafmid BA; Site 1: HindIII;  
Site 2: NotI; sex=Female; dev stage=3 months old;  
isolate=muscular atrophy patient; tissue\_type=total brain  
Total mRNA was oligo-(dT) primed and directionally  
cloned 5' -> 3' into the HindIII -> NotI sites of the  
lafmid BA vector. Clone library from B.Souares, Psychiatry  
Dept. Columbia University, USA. Normalization\_method:  
Bento Soares, P.N.A.S in press"

BASE COUNT  
ORIGIN  
88 a 42 c 43 g 69 t

Query Match 80.0%; Score 16; DB 14; Length 242;  
Best Local Similarity 100.0%; Pred.No. 68;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 2 CTTAGGAGACCACT 17  
|||||  
48 CTTAGGAGACCACT 63

RESULT 6  
LOCUS AU231688 350 bp mRNA linear EST 21-SEP-2001  
DEFINITION AU231688 Cloned bovine fetus cDNA Bos taurus clone Cln595 3',  
mRNA sequence.

ACCESSION AU231688  
VERSION AU231688.1 GI:15719980  
KEYWORDS EST.  
SOURCE Bos taurus (cow)  
ORGANISM Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;  
Bovidae; Bovinae; Bos.

REFERENCE  
AUTHORS  
TITLE  
JOURNAL  
COMMENT  
1 (bases 1 to 350)  
Taniguchi, Y., Lejokole, H.Y., Yamada, T., Akagi, S., Yasue, H. and  
Sasaki, Y.  
Analysis of expressed sequence tags from a cDNA library of somatic  
nuclear transfer-derived cloned bovine fetus  
Unpublished  
Contact: Takahisa Yamada  
Graduate School of Agriculture

Kyoto University  
Saiyoku, Kitashirakawa, Kyoto, Kyoto 606-8502, Japan  
Tel: 81-75-753-6323  
Fax: 81-75-753-6340  
Email: tyamada@kane.kans.kais.kyoto-u.ac.j  
This clone was obtained from a 3' end cDNA library.  
Location/Qualifiers

FEATURES  
source

1..350  
/organism="Bos taurus"  
/mol\_type="mRNA"  
/db\_xref="taxon:9913"  
/clone="Cln595"  
/dev\_stage="fetus"  
/clone\_lib="Cloned bovine fetus cDNA"  
BASE COUNT 101 a 70 c 83 g 92 t 4 others  
ORIGIN

Query Match 80.0%; Score 16; DB 9; Length 350;  
Best Local Similarity 100.0%; Pred.No. 73;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 2 CTTAGGAGACCACT 17  
|||||  
252 CTTAGGAGACCACT 267

RESULT 7  
LOCUS R24852 358 bp mRNA linear EST 20-APR-1995  
DEFINITION y929605.r1 Soares infant brain INIB Homo sapiens cDNA clone  
IMAGE:33804 5', mRNA sequence.

ACCESSION R24852  
VERSION R24852.1 GI:779740  
KEYWORDS EST.  
SOURCE Homo sapiens (human)  
ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.  
1 (bases 1 to 358)

REFERENCE  
AUTHORS  
TITLE  
JOURNAL  
COMMENT  
Hillier, L., Clark, N., Dubuque, T., Elliston, K., Hawkins, M., Holman, R.,  
Rifkin, L., Rohlfing, T., Soares, M., Tan, F., Tivadaris, E., Waterston, R.,  
Williamson, A., Wohlmann, P. and Wilson, R.  
The Washu-Merck EST Project  
Unpublished  
Contact: Wilson R.K  
Washington University School of Medicine  
444 Forest Park Parkway, Box 8501, St. Louis, MO 63108  
Tel: 314 286 1800  
Fax: 314 286 1810  
Email: est@watson.wustl.edu  
Insert Size: 2215  
High quality sequence stops: 284 Source: IMAGE Consortium, LNLN.  
This clone is available royalty-free through LNLN; contact the  
IMAGE Consortium (info@image.lnl.gov) for further information.  
Insert Length: 2215 Std Error: 0.00  
Seq primer: M13RPI  
High quality sequence stop: 284.  
Location/Qualifiers

FEATURES  
source

1..358  
/organism="Homo sapiens"  
/mol\_type="mRNA"  
/db\_xref="GDB:406151"  
/db\_xref="taxon:9606"  
/clone="IMAGE:33804"  
/sex="Female"  
/dev\_stage="73 days post natal"  
/lab\_host="DH10B (ampicillin resistant)"  
/clone\_lib="Soares infant brain INIB"  
/note="Organ: whole brain; Vector: lafmid BA; Site 1: Not  
I; Site 2: Hind III; 1st strand cDNA was primed with a Not  
I - oligo(dT) primer [5'  
ACTGGAAGATTCGGCGCGAGGAAATTTTATTTTATTTT 3'];

double-stranded cDNA was ligated to Hind III adaptors (Pharmacia), digested with Not I and directionally cloned into the Not I and Hind III sites of the lacMid BA vector. Library went through one round of normalization. Library constructed by Bento Soares and M. Fatima Bonaldo."

BASE COUNT 112 a 64 c 68 g 112 t 2 others  
 ORIGIN

Query Match 80.0%; Score 16; DB 14; Length 358;  
 Best Local Similarity 100.0%; Pred. No. 74;  
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 CTTAGAGGACAACT 17  
 47 CTTAGAGGACAACT 62

RESULT 8  
 LOCUS AQ483201 390 bp DNA linear GSS 24-APR-1999  
 DEFINITION RPCI-11-241P16.TV RPCI-11 Homo sapiens genomic clone RPCI-11-241P16  
 ACCESSION AQ483201  
 VERSION AQ483201.1 GI:4670605  
 KEYWORDS GSS.  
 SOURCE Homo sapiens (human)  
 ORGANISM Homo sapiens

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo. 1 (bases 1 to 390)  
 AUTHORS Zhao, S., Adams, M.D., Nierman, W., Malek, J., de Jong, P. and Venter, J.C.  
 TITLE Use of BAC End Sequences from Library RPCI-11 for Sequence-Ready Map Building  
 JOURNAL Unpublished  
 COMMENT Other GSSs: RPCI-11-241P16.TV  
 Contact: Shaying Zhao, William Nierman, Mark Adams  
 Department of Eukaryotic Genomics  
 The Institute for Genomic Research  
 9712 Medical Center Dr., Rockville, MD 20850  
 Tel: 301 838 0200  
 Fax: 301 838 0208  
 Email: hbe@tigr.org  
 Clones are derived from the human BAC library RPCI-11. For BAC library availability, please contact Pieter de Jong (pde@tigr.org) or from BACPAC Resources (http://bacpac.med.buffalo.edu/ordering) or from Research Genet. cs (info@resgen.com). BAC end search page: http://www.tigr.org/tdb/hunguen/bac\_end\_search/bac\_end\_search.html.  
 Seg primer: SP6  
 Classes: BAC ends.

FEATURES  
 source  
 location/Qualifiers

1..390  
 /organism="Homo sapiens"  
 /mol\_type="genomic DNA"  
 /db\_xref="GBB:7592535"  
 /db\_xref="taxon:9606"  
 /clone="RPCI-11-241P16"  
 /sex="Male"  
 /cell\_type="Tymphocytes"  
 /clone\_lib="RPCI-11"  
 /note="Vector: pBACe3.6; Site 1: EcoRI; Site 2: EcoRI; RPCI11 Human Male BAC Library"  
 BASE COUNT 112 a 85 c 89 g 104 t  
 ORIGIN

Query Match 80.0%; Score 16; DB 28; Length 390;  
 Best Local Similarity 100.0%; Pred. No. 75;  
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 AGGAGGACAACTCCC 20  
 289 AGGAGGACAACTCCC 304  
 DB

RESULT 9  
 LOCUS A1696726 395 bp mRNA linear EST 17-DEC-1999  
 DEFINITION wc56d11.x1 NCI CGAP Pr28 Homo sapiens cDNA clone IMAGE:232645 3' similar to contains ORF.t1 ORF ORF repetitive element ;, mRNA sequence.

ACCESSION A1696726  
 VERSION A1696726.1 GI:4984626  
 KEYWORDS EST.  
 SOURCE Homo sapiens (human)  
 ORGANISM Homo sapiens

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo. 1 (bases 1 to 395)  
 AUTHORS NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.  
 TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP), Tumor Gene Index  
 JOURNAL Unpublished  
 COMMENT Contact: Robert Strausberg, Ph.D.  
 Email: cgapbs-remail.nih.gov  
 Tissue Procurement: Michael J. Brownstein, M.D., Ph.D., Michael R. Emmert-Buck, M.D., Ph.D.  
 CDNA Library Preparation: M. Bento Soares, Ph.D.  
 DNA Sequencing by: Washington University Genome Sequencing Center  
 Clone distribution: NCI-CGAP clone distribution information can be found through the I.M.A.G.E. Consortium/LNL at: www-bio.llnl.gov/bdip/image/image.html  
 Insert Length: 551 Std Error: 0.00  
 Seg primer: -40UP from Gibco  
 High quality sequence stop: 394.

FEATURES  
 source  
 location/Qualifiers

1..395  
 /organism="Homo sapiens"  
 /mol\_type="mRNA"  
 /db\_xref="taxon:9606"  
 /clone="IMAGE:232645"  
 /sex="male"  
 /dev\_stage="adult"  
 /lab\_host="DH10B"  
 /clone\_lib="NCI CGAP Pr28"  
 /note="Organ: prostate; Vector: pT73D-Pac (Pharmacia) with a modified polylinker. Plasmid DNA from the normalized library NCI CGAP Pr22 was prepared, and 85 clones were made in vitro. Following HAP purification, this DNA was used as tracer in a subtractive hybridization reaction. The driver was PCR-amplified cDNAs from a pool of 5,000 clones made from the same library (clones 985608-986759, 1101192-1101959, and 1217928-1220615). Subtraction by Bento Soares and M. Fatima Bonaldo."  
 BASE COUNT 125 a 85 c 97 g 88 t  
 ORIGIN

Query Match 80.0%; Score 16; DB 9; Length 395;  
 Best Local Similarity 100.0%; Pred. No. 75;  
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 AGGAGGACAACTCCC 20  
 84 AGGAGGACAACTCCC 99  
 DB

RESULT 10  
 LOCUS T75305 400 bp mRNA linear EST 03-MAR-1995  
 DEFINITION YC89C08.r1 Soares infant brain INIB Homo sapiens cDNA clone IMAGE:23341 5', mRNA sequence.  
 ACCESSION T75305  
 VERSION T75305.1 GI:692067  
 KEYWORDS EST.  
 SOURCE Homo sapiens (human)

ORGANISM Homo sapiens  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.  
 REFERENCE 1 (bases 1 to 400)  
 AUTHORS Hillier, L., Clark, N., Dubuque, T., Elliston, K., Hawkins, M., Holman, M., Hultman, M., Kucaba, T., Le, M., Lennon, G., Marra, M., Parsons, J., Rifkin, B., Rohlfing, T., Soares, M., Tan, F., Trevisan, E., Waterston, R., Williamson, A., Woldmann, P. and Wilson, R.  
 TITLE The Washu-Merck EST Project  
 JOURNAL Unpublished  
 COMMENT Contact: Wilson RK  
 Washington University School of Medicine  
 444 Forest Park Parkway, Box 8501, St. Louis, MO 63108  
 Tel: 314 286 1800  
 Fax: 314 286 1810  
 Email: est@wustl.edu  
 Insert Size: 2260  
 High quality sequence stops: 318 Source: IMAGE Consortium, LNL  
 This clone is available royalty-free through LNL; contact the IMAGE Consortium (info@image.llnl.gov) for further information.  
 Insert Length: 2260 Std Error: 0.00  
 Seq primer: M3RP1  
 High quality sequence stop: 318.  
 Location/Qualifiers  
 1..400  
 /organism="Homo sapiens"  
 /mol\_type="mRNA"  
 /db\_xref="GDB:395688"  
 /db\_xref="taxon:9606"  
 /clone="IMAGE:23341"  
 /sex="female"  
 /dev\_stage="73 days post natal"  
 /lab\_host="DH10B (ampicillin resistant)"  
 /clone\_lib="Soares Infant Brain INB"  
 /note="Organ: whole brain; Vector: Latmid BA; Site: 1: Not I; Site 2: Hind III; 1st strand cDNA was primed with a Not I - oligo(dT) primer [5', AACTGAGAGAAATCGCGCCGACGAGAAATTTTTTTTTTTTTTTT 3']; double-stranded cDNA was ligated to Hind III adaptors (Pharmacia), digested with Not I and directionally cloned into the Not I and Hind III sites of the Latmid BA vector. Library went through one round of normalization. Library constructed by Bento Soares and M. Patricia Bonaldo."

BASE COUNT 123 a 77 c 72 g 127 t 1 others

ORIGIN

Query Match 80.0%; Score 16; DB 14; Length 400;  
 Best Local Similarity 100.0%; Pred. No. 75;  
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 2 CTTAGAGAGAACACT 17  
 ||||||||||||  
 Db 26 CTTAGAGAGAACACT 41

RESULT 11  
 LOCUS AQ379270 413 bp DNA linear GSS 20-MAY-1999  
 DEFINITION RPI11-151H4.TJ RPI1-11 Homo sapiens genomic clone RPI1-11-151H4, genomic survey sequence.  
 ACCESSION AQ379270  
 VERSION AQ379270.1 GI:4350293  
 KEYWORDS GSS.  
 SOURCE Homo sapiens (human)  
 ORGANISM Homo sapiens  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.  
 REFERENCE 1 (bases 1 to 413)  
 AUTHORS Zhao, S., Adams, M. D., Nieman, W., Malek, J., de Jong, P. and Venter, J. C.  
 TITLE Use of BAC End Sequences from Library RPI1-11 for Sequence-Ready Map Building  
 JOURNAL Unpublished

COMMENT Other GSSs: RPI11-151H4.TJ  
 Contact: Shaying Zhao, William Nieman, Mark Adams  
 Department of Eukaryotic Genomics  
 The Institute for Genomic Research  
 9712 Medical Center Dr., Rockville, MD 20850  
 Tel: 301 838 0200  
 Fax: 301 838 0208  
 Email: hbe@tigr.org  
 Clones are derived from the human BAC library RPI1-11. For BAC library availability, please contact Pieter de Jong (pieter@dejong.med.buffalo.edu). Clones may be purchased from BACPAC Resources (http://bacpac.med.buffalo.edu/ordering) or from Research Genetics (info@resgen.com). BAC end search page: http://www.tigr.org/tdb/humgen/bac\_end\_search/bac\_end\_search.html  
 Seq primer: SP6  
 Class: BAC ends.  
 Location/Qualifiers  
 1..413  
 /organism="Homo sapiens"  
 /mol\_type="genomic DNA"  
 /db\_xref="GDB:7557771"  
 /db\_xref="taxon:9606"  
 /clone="RPI1-11-151H4"  
 /sex="male"  
 /cell\_type="Lymphocytes"  
 /clone\_lib="RPI1-11"  
 /note="Vector: pBAC3.6; Site 1: EcoRI; Site 2: EcoRI; RPI11 Human Male BAC library"

BASE COUNT 124 a 92 c 92 g 104 t 1 others

ORIGIN

Query Match 80.0%; Score 16; DB 28; Length 413;  
 Best Local Similarity 100.0%; Pred. No. 76;  
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 5 AGAGAGAACAGTCCC 20  
 ||||||||||||  
 Db 273 AGAGAGAACAGTCCC 288

RESULT 12  
 LOCUS AM769407 431 bp mRNA linear EST 04-MAY-2000  
 DEFINITION h163g12.x1 NCI CGAP Kid13 Homo sapiens cDNA clone IMAGE:3005926 3' similar to confins\_OPR.b1 OPR repetitive element ;, mRNA sequence.  
 ACCESSION AM769407  
 VERSION AM769407.1 GI:7701438  
 KEYWORDS EST.  
 SOURCE Homo sapiens (human)  
 ORGANISM Homo sapiens  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.  
 REFERENCE 1 (bases 1 to 431)  
 AUTHORS NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.  
 TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP), Tumor Gene Index  
 JOURNAL Unpublished  
 COMMENT Contact: Robert Strausberg, Ph.D.  
 Email: cgapbs-remail.nih.gov  
 Tissue Procurement: Chris Moskaluk, M.D., Ph.D., Michael R. Emmert-Buck, M.D., Ph.D. CDNA Library Preparation: Life Technologies, Inc. CDNA Library Arrayed by: Christa Prange, The I.M.A.G.E. Consortium DNA Sequencing by: Washington University Genome Sequencing Center  
 Clone distribution: NCI-CGAP clone distribution information can be found through the I.M.A.G.E. Consortium/LNL, send email to: info@image.llnl.gov  
 Possible reversed clone: polyT not found  
 Seq primer: -40up from Gibco  
 High quality sequence stop: 396.  
 Location/Qualifiers  
 1..431  
 /organism="Homo sapiens"

FEATURES  
 source

/mol\_type="mRNA"  
/db\_xref="taxon:9606"  
/clone="IMAGE:3005926"  
/tissue\_type="2 pooled Wilms' tumors, one primary and one metastatic to brain"  
/lab\_host="DH10B"  
/clone\_1fb="NCI CGAP Kid13"  
/note="Organ: kidney; Vector: PCMV-SPORE6; Site 1: SalI; Site 2: NotI; Cloned unidirectionally. Primer: Oligo dt. library constructed by Life Technologies."  
Library constructed by Life Technologies."

BASE COUNT 131 a 98 c 103 g 99 t

Query Match 80.0%; Score 16; DB 9; Length 431;  
Best Local Similarity 100.0%; Pred. No. 77;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 5 AGGAGGAACAAGTCCC 20  
89 AGGAGGAACAAGTCCC 104

RESULT 13  
BE227333 443 bp mRNA linear EST 06-JUL-2000  
LOCUS 894030E10.x3 C. reinhardtii CC-1690, normalized, lambda zap II  
DEFINITION Chlamydomonas reinhardtii cDNA, mRNA sequence.  
ACCESSION BE227333  
VERSION BE227333.1 GI:8932572  
KEYWORDS EST.  
SOURCE Chlamydomonas reinhardtii  
ORGANISM Chlamydomonas reinhardtii  
Eukaryota; Viridiplantae; Chlorophyta; Chlorophyceae; Volvocales;  
Chlamydomonadaceae; Chlamydomonas.  
1 (bases 1 to 443)  
Grossman, A., Davies, J., Federspiel, N., Harris, E., Lefebvre, P., McDermott, J. P., Silflow, C., Stern, D. and Surzycki, R.  
Analyses of the Chlamydomonas reinhardtii Genome: A Model, Unicellular System for Analyzing Gene Function and Regulation in Vascular Plants; project phase 2  
Unpublished  
Contact: Elizabeth H. Harris  
DCMB Box 91000  
Duke University  
Durham, NC 27708-1000, USA  
Tel: 919 613 8164  
Fax: 919 613 8177  
Email: chlamy@duke.edu.  
Location/Qualifiers  
1. 443  
/organism="Chlamydomonas reinhardtii"  
/mol\_type="mRNA"  
/strain="CC-1690 wild type mt+ 21gr"  
/db\_xref="taxon:3055"  
/clone\_1fb="C. reinhardtii CC-1690, normalized, lambda zap II"  
/note="Vector: pBluescript II SK-; Site 1: EcoRI; Site 2: XhoI; This library, constructed by John Davies and Jeffrey McDermott, combines cDNAs from CC-1690 cells grown to mid-log phase in TAP (acetate-containing) medium in the light, TAP medium in the dark, HS (minimal) medium in ambient levels of CO2 and HS medium bubbled with 5% CO2. PolyA mRNA was purified from each sample, pooled and cDNA synthesized. The cDNA was directionally cloned into lambda ZAP II (Stratagene) in the EcoRI (5') and XhoI (3') sites. pBluescript II SK- plasmids were excised from the lambda ZAP clones by superinfection with Exsistat (Stratagene) phage. The library was normalized using method 4 described in Bonaldo et al (1996) Genome Research 6: 791-806."

BASE COUNT 89 a 123 c 139 g 86 t

Query Match 80.0%; Score 16; DB 10; Length 443;

Best Local Similarity 100.0%; Pred. No. 77;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 5 AGGAGGAACAAGTCCC 20  
29 AGGAGGAACAAGTCCC 44

RESULT 14  
BY414889 464 bp mRNA linear EST 13-DEC-2002  
LOCUS BY414889 RIKEN full-length enriched, 16 days embryo kidney Mus  
DEFINITION musculus cDNA clone 1920006j23 3', mRNA sequence.  
ACCESSION BY414889  
VERSION BY414889.1 GI:26679833  
KEYWORDS EST.  
SOURCE Mus musculus (house mouse)  
ORGANISM Mus musculus  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mus.  
1 (bases 1 to 464)  
Okazaki, Y., Furuno, M., Kasukawa, T., Adachi, J., Bono, H., Kondo, S., Nikaido, I., Osato, N., Sato, R., Suzuki, H., Yamanka, I., Kiyosawa, H., Yagi, K., Tomaru, Y., Hasegawa, Y., Nogami, A., Schonbach, C., Gajdosi, T., Baldarelli, R., Hill, D. P., Butt, C., Hume, D. A., Quackenbush, J., Schriml, L. M., Kanapin, A., Matsuda, H., Batilov, S., Bisset, K. W., Blake, J. A., Bradt, P., Brusic, V., Choitha, C., Forrest, L. E., Cousins, S., Dalla, E., Dragan, T. A., Fletcher, C. F., Forrest, A., Fraser, K. S., Gaasterland, T., Gariboldi, M., Gissi, C., Godzik, A., Gough, J., Grimmond, S., Gustincich, S., Hirokawa, N., Jackson, J. J., Jarvis, E. D., Kanai, A., Kawai, H., Kawasawa, Y., Kedzierski, R. M., King, B. L., Konagaya, A., Kurochkin, I. V., Lee, Y., Lenhard, B., Lyons, P. A., Maglott, D. R., Malais, L., Marchionni, L., McKenzie, L., Miki, H., Nagashima, T., Numata, K., Okido, T., Pavan, W. J., Pereira, G., Pesole, G., Petrovsky, N., Pillai, R., Pontius, J. U., Qi, D., Ramachandran, S., Ravasi, T., Reed, J. C., Reed, D. J., Reid, J., Ring, B. Z., Ringwald, M., Sanderlin, A., Schneider, C., Semple, C. A., Setou, M., Shimada, K., Sultana, R., Takenaka, Y., Taylor, M. S., Teasdale, R. D., Tomita, M., Verardo, R., Wagner, L., Wahlestedt, C., Wang, Y., Watanabe, Y., Wells, C., Wilming, L. G., Wyshaw-Boris, A., Yangisawa, M., Yang, L., Yang, L., Yuan, Z., Zavolan, M., Zhu, Y., Zimmer, A., Carninci, P., Hayatsu, N., Hirozane-Kishikawa, T., Kono, H., Nakamura, M., Sakazume, N., Sato, K., Shiraki, T., Waki, K., Kawai, J., Aizawa, K., Arakawa, T., Fukuda, S., Hara, A., Hashizume, W., Imotani, K., Ishii, Y., Itoh, M., Kagawa, I., Miyazaki, A., Sakai, K., Sasaki, D., Shibata, K., Shinagawa, A., Yasunishi, A., Yoshino, M., Waterston, R., Lander, E. S., Rogers, J., Birney, E. and Hayashizaki, Y.  
Analysis of the mouse transcriptome based on functional annotation of 60,770 full-length cDNAs  
Nature 420, 563-573 (2002)  
22354683  
MEDLINE  
PUBMED 12466851

JOURNAL COMMENT  
TITLE  
CONTACT: Yoshihide Hayashizaki  
Laboratory for Genome Exploration Research Group, RIKEN Genomic Sciences Center (GSC), Yokohama Institute  
The Institute of Physical and Chemical Research (RIKEN)  
1-7-22 Suenhiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan  
Tel: 81-45-503-9222  
Fax: 81-45-503-9216  
Email: genome-resgsc.riken-go.jp/  
URL: http://genome-gsc.riken-go.jp/  
Aizawa, K., Akimura, T., Arakawa, T., Carninci, P., Fukuda, S., Hirozane, T., Imotani, K., Ishii, Y., Itoh, M., Kawai, J., Kono, H., Miyazaki, A., Murata, M., Nakamura, M., Nomura, K., Numazaki, R., Ohno, M., Sakai, K., Sakazume, N., Sasaki, D., Sato, K., Shibata, K., Shiraki, T., Tagami, M., Waki, K., Watanabe, A., Watanabe, M. and Hayashizaki, Y. Direct Submission  
Computational Analysis of Full-length Mouse cDNAs Compared with Human Genome Sequences Mamm. Genome 12, 673-677 (2001)  
Normalization and subtraction of cap-trapper-selected cDNAs to prepare full-length cDNA libraries for rapid discovery of new genes. Genome Res. 10 (10), 1617-1630 (2000)  
RIKEN integrated sequence analysis (RISA) system-384-format

sequencing pipeline with 384 multicapillary sequencer. Genome Res. 10 (11), 1757-1771 (2000)  
 Computer-based methods for the mouse full-length cDNA encyclopedia: Real-time sequence clustering for construction of a nonredundant cDNA library. Genome Res. 11 (2), 281-289 (2001)  
 cDNA library was prepared and sequenced in Mouse Genome Encyclopedia Project of Genome Exploration Research Group in Riken Genomic Sciences Center and Genome Science Laboratory in RIKEN Division of Experimental Animal Research in Riken contributed to prepare mouse tissues.  
 Please visit our web site (<http://genome.gsc.riken.go.jp>) for further details.

## FEATURES

## source

Location/Qualifiers

1..464

/organism="Mus musculus"

/mol\_type="mRNA"

/strain="C57BL/6J"

/db\_xref="taxon:10090"

/clone="192006J23"

/tissue\_type="kidney"

/dev\_stage="16 days embryo"

/clone\_lib="RIKEN full-length enriched, 16 days embryo kidney"

BASE COUNT 132 a 84 c 101 g 146 t 1 others

## ORIGIN

Query Match 80.0%; Score 16; DB 13; Length 464;  
 Best Local Similarity 100.0%; Pred. No. 78;  
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 AGAGAGAACAAAGTCCC 20

Db 241 AGAGAGAACAAAGTCCC 256

## RESULT 15

## LOCUS

R24326 476 bp mRNA linear EST 20-APR-1995

DEFINITION Y932f03.r1 Soares infant brain JN1B Homo sapiens cDNA clone

IMAGE:33987 5', mRNA sequence.

VERSION R24326.1 GI:779214

KEYWORDS EST.

SOURCE Homo sapiens (human)

ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.

REFERENCE 1 (bases 1 to 476)

Hillier, L., Clark, N., Dubuque, T., Elliston, K., Hawkins, M., Holman

, M., Hultman, M., Kucaba, T., Le, M., Lennon, G., Marra, M., Parsons, J.,

Rifkin, L., Rohlfing, T., Soares, M., Tan, F., Trevaskis, E., Waterston

, R., Williamson, A., Woldmann, P. and Wilson, R.

The Washu-Merck EST Project

Unpublished

JOURNAL

COMMENT

Washington University School of Medicine

4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108

Tel: 314 286 1800

Fax: 314 286 1810

Email: [est@watson.wustl.edu](mailto:est@watson.wustl.edu)

High quality sequence strops: 349

Source: IMAGE Consortium, LNL

This clone is available royalty-free through LNL; contact the

IMAGE Consortium ([info@image.llnl.gov](mailto:info@image.llnl.gov)) for further information.

Seq primer: M13P1

High quality sequence strop: 349.

## FEATURES

## source

1..476

/organism="Homo sapiens"

/mol\_type="mRNA"

/db\_xref="GDB:406334"

/db\_xref="taxon:9606"

/clone="IMAGE:33987"

/sex="female"  
 /dev\_stage="73 days post natal"  
 /lab\_host="DH10B (ampicillin resistant)"  
 /clone\_lib="Soares infant brain JN1B"  
 /note="Organ: whole brain; Vector: lambda B; Site 1: Not  
 I; Site 2: Hind III; 1st strand cDNA was primed with a Not  
 I - oligo(dT) primer [5]  
 ACTGAGAGAAATTCGCGCGCGAGATTTTCTTTTCTTTT 3'1];  
 double-stranded cDNA was ligated to Hind III adaptors  
 (Pharmacia), digested with Not I and directionally cloned  
 into the Not I and Hind III sites of the lambda B vector.  
 Library went through one round of normalization. Library  
 constructed by Bento Soares and M. Fatima Bonaldo."

BASE COUNT 154 a 87 c 86 g 148 t 1 others

## ORIGIN

Query Match 80.0%; Score 16; DB 14; Length 476;  
 Best Local Similarity 100.0%; Pred. No. 78;  
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 CTTAGAGAGAACAAAGT 17

Db 47 CTTAGAGAGAACAAAGT 62

## RESULT 16

## LOCUS

BF830043/c 487 bp mRNA linear EST 13-JAN-2001

DEFINITION MR3-HN0052-271200-001-b01 HN0052 Homo sapiens cDNA, mRNA sequence.

VERSION BF830043.1 GI:12176257

KEYWORDS EST.

SOURCE Homo sapiens (human)

ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.

REFERENCE 1 (bases 1 to 487)

Dias Neto, E., Garcia Correa, R., Verjovski-Almeida, S., Briones, M.R.,

Nagai, M.A., da Silva, W. Jr., Zago, M.A., Bordin, S., Costa, F.F.,

Goldman, G.H., Carvalho, A.F., Matsukuma, A., Bata, G.S., Simpson, D.H.,

Brunstein, A., de Oliveira, P.S., Bucher, P., Jongeneel, C.V., O'Hare

, M.J., Soares, F., Brenhant, R.R., Reis, L.F., de Souza, S.J. and

Simpson, A.J.

Shotgun sequencing of the human transcriptome with ORF expressed

sequence tags

Proc. Natl. Acad. Sci. U.S.A. 97 (7), 3491-3496 (2000)

2020263

10737800

Contact: Simpson A.J.G.

Laboratory of Cancer Genetics

Ludwig Institute for Cancer Research

Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP,

Brazil

Tel: +55-11-2704922

Fax: +55-11-2707001

Email: [asimpson@ludwig.org.br](mailto:asimpson@ludwig.org.br)

This sequence was derived from the FAPSP/LICR Human Cancer Genome

Project. This entry can be seen in the following URL

(<http://www.ludwig.org.br/scripts/gethtml2.pl?l=MR3&t2=MR3-HN0052-271200-001-b01&t3=2000-12-27&t4=1>)

Seq primer: puc 18 forward

High quality sequence start: 32

High quality sequence stop: 463.

Location/Qualifiers

1..487

/organism="Homo sapiens"

/mol\_type="mRNA"

/db\_xref="taxon:9606"

/dev\_stage="Adult"

/clone\_lib="HN0052"

/note="Organ: head normal; Vector: puc18; Site 1: SmaI;

Site 2: SmaI; A mini-library was made by cloning products

derived from ORESTES PCR (U.S. letters patent application

No. 196,716 - Ludwig Institute for Cancer Research)  
profiles into the pUC 18 vector. Reverse transcription of  
tissue mRNA and cDNA amplification were performed under  
low stringency conditions."

BASE COUNT 165 a 102 c 108 g 112 t  
ORIGIN

Query Match 80.0%; Score 16; DB 10; Length 487;  
Best Local Similarity 100.0%; Pred. No. 79;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 2 CTTAGAGGAGCAACAGT 17  
483 CTTAGAGGAGCAACAGT 468

RESULT 17  
R24340 494 bp mRNA linear EST 20-APR-1995  
LOCUS YG32h03.r1 Soares infant brain INIB Homo sapiens cDNA clone  
DEFINITION IMAGE:34200 5', mRNA sequence.

ACCESSION R24340  
VERSION R24340.1 GI:779228  
KEYWORDS EST.  
SOURCE Homo sapiens (human)  
ORGANISM Homo sapiens

REFERENCE 1 (bases 1 to 494)  
AUTHORS Hillier, L., Clark, N., Dubuque, T., Elliston, K., Hawkins, M., Holman, M., Hultman, M., Kucaba, T., Le, M., Lennon, G., Marra, M., Parsons, J., Rikkin, L., Rohlfing, T., Soares, M., Tan, F., Trevisakis, E., Waterston, R., Williamson, A., Wohlmann, P. and Wilson, R.  
TITLE The WashU-Merck EST Project  
JOURNAL Unpublished  
COMMENT Contact: Wilson RK  
Washington University School of Medicine  
444 Forest Park Parkway, Box 8501, St. Louis, MO 63108  
Tel: 314 286 1800  
Fax: 314 286 1810  
Email: eest@watson.wustl.edu  
Insert Size: 2185  
High quality sequence stops: 325 Source: IMAGE Consortium, LLNL  
This clone is available royalty-free through LLNL; contact the  
IMAGE Consortium (info@image.llnl.gov) for further information.  
Insert Length: 2185 Std Error: 0.00  
Seq primer: M13RP1  
High quality sequence stop: 325.

#### FEATURES

source

1: 494  
/organism="Homo sapiens"  
/mol\_type="mRNA"  
/db\_xref="GDB:406547"  
/db\_xref="taxon:9606"  
/clone="IMAGE:34200"  
/sex="female"  
/dev\_stage="73 days post natal"  
/lab\_host="DH10B (ampicillin resistant)"  
/clone\_1lb="Soares infant brain INIB"  
/note="Organ: whole brain; Vector: Latmid B; Site: 1: Not  
I; Site 2: Hind III; 1st strand cDNA was primed with a Not  
I - oligo (dT) primer [5',  
AAGTGAAGATTCGCGCCGAGCAATTTTCTTTTCTTTT 3'];  
double-stranded cDNA was ligated to Hind III adaptors  
(Pharmacia), digested with Not I and directionally cloned  
into the Not I and Hind III sites of the Latmid B vector.  
Library went through one round of normalization. Library  
constructed by Bento Soares and M. Patricia Bonaldo."

BASE COUNT 151 a 90 c 90 g 159 t 4 others  
ORIGIN

Query Match 80.0%; Score 16; DB 14; Length 494;  
Best Local Similarity 100.0%; Pred. No. 79;

Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
OY 2 CTTAGAGGAGCAACAGT 17  
47 CTTAGAGGAGCAACAGT 62  
DB

RESULT 18  
BB693705/c 516 bp mRNA linear EST 10-OCT-2001  
LOCUS BB693705 RIKEN full-length enriched, 2 days neonate sympathetic  
ganglion Mus musculus cDNA clone 7120448G11 3', mRNA sequence.  
DEFINITION BB693705  
ACCESSION BB693705  
VERSION BB693705.1 GI:16020438  
KEYWORDS EST.  
SOURCE Mus musculus (house mouse)  
ORGANISM Mus musculus

REFERENCE 1 (bases 1 to 516)  
AUTHORS Akimura, T., Arakawa, T., Carninci, P., Furuno, M., Hangaki, T., Hayatsu, N., Hiramoto, K., Hirooka, T., Hirozane, T., Imotani, K., Ishii, Y., Ito, M., Kawai, J., Kojima, Y., Kono, H., Kouda, M., Matsuyama, T., Nakamura, M., Nishi, K., Nomura, K., Numasaki, R., Okazaki, Y., Okido, T., Saito, R., Sakai, C., Sakai, K., Sakazume, N., Sasaki, D., Sato, K., Shibata, K., Shinagawa, A., Shiraki, T., Sogabe, Y., Suzuki, H., Tagawa, A., Takahashi, F., Takaku-Akita, S., Tanaka, T., Tomaru, A., Toyota, T., Watabiki, A., Yasunishi, A., Muramatsu, M. and Hayashizaki, Y.  
TITLE RIKEN Encyclopedia of Mouse Full-length cDNAs (Akimura, T., et al. 2001)  
JOURNAL Unpublished  
COMMENT Contact: Yoshitake Hayashizaki  
Laboratory for Genome Exploration Research Group, RIKEN Genomic  
Sciences Center (GSC), Yokohama Institute  
The Institute of Physical and Chemical Research (RIKEN)  
1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan  
Tel: 81-45-503-9222  
Fax: 81-45-503-9216  
Email: genome-res@gsr.riken.go.jp,  
URL: http://genome.gsc.riken.go.jp/  
Carninci, P., Shibata, Y., Hayatsu, N., Sugahara, Y., Shibata, K., Itoh, M., Kono, H., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y.  
Normalization and subtraction of cap-trapper-selected cDNAs to  
prepare full-length cDNA libraries for rapid discovery of new  
genes. Genome Res. 10 (10), 1617-1630 (2000)  
wagi, K., Fujiwara, S., Inoue, K., Togawa, Y., Izawa, M., Ohara, E., Watabiki, M., Yoneda, Y., Ishikawa, T., Ozawa, K., Tanaka, T., Matsura, S., Kawai, J., Okazaki, Y., Muramatsu, M., Inoue, Y., Kira, A. and Hayashizaki, Y.  
RIKEN integrated sequence analysis (RISA) system--384-format  
sequencing pipeline with 384 multicapillary sequencer. Genome Res. 10 (11), 1757-1771 (2000)  
Kono, H., Fukunishi, Y., Shibata, K., Itoh, M., Carninci, P., Sugahara, Y. and Hayashizaki, Y.  
Computer-based method for the mouse full-length cDNA  
encyclopedia: real-time sequence clustering for construction of a  
nonredundant cDNA library. Genome Res. 11 (2), 281-289 (2001)  
Please visit our web site (http://genome.gsc.riken.go.jp) for  
further details.  
e mouse tissues.

#### FEATURES

source

1: 516  
/organism="Mus musculus"  
/mol\_type="mRNA"  
/strain="C57BL/6J"  
/db\_xref="taxon:10090"  
/clone="7120448G11"  
/sex="mixed"  
/tissue\_type="sympathetic ganglion"  
/dev\_stage="2 days neonate"  
/lab\_host="DH10B"  
/clone\_1lb="RIKEN full-length enriched, 2 days neonate  
sympathetic ganglion"







LOCUS B0347567 550 bp mRNA linear EST 20-MAY-2002  
 DEFINITION CM0-HT0179-051099-064-g10 HT0179 Homo sapiens cDNA, mRNA sequence.  
 ACCESSION B0347567  
 VERSION B0347567.1 GI:21011623  
 KEYWORDS EST.  
 SOURCE Homo sapiens (human)  
 ORGANISM Homo sapiens  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.  
 1 (bases 1 to 550)  
 Dias Neto,E., Garcia Correa,R., Verjovski-Almeida,S., Briomes,M.R., Nagai,M.A., da Silva,W.U., Zago,M.A., Bordin,S., Costa,F.F., Goldman,G.H., Carvalho,A.F., Matukuma,A., Bala,G.S., Simpson,D.H., Brunstein,A., deOliveira,P.S., Bucher,P., Jongeneel,C.V., O'Hare,M.J., Soares,F., Brentani,R.R., Reis,L.F., de Souza,S.J. and Simpson,A.J.  
 Shotgun sequencing of the human transcriptome with ORF expressed sequence tags  
 Proc. Natl. Acad. Sci. U.S.A. 97 (7), 3491-3496 (2000)  
 2020263  
 MEDLINE 10737800  
 PUBMED  
 COMMENT Contact: Simpson A.J.G.  
 Laboratory of Cancer Genetics  
 Ludwig Institute for Cancer Research  
 Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP, Brazil  
 Tel: +55-11-2704922  
 Fax: +55-11-2707001  
 Email: asimpso@ludwig.org.br  
 This sequence was derived from the PAPESP/LICR Human Cancer Genome Project. This entry can be seen in the following URL  
 (http://www.ludwig.org.br/scripts/gethtml2.pl?tl=CM0&lt2=CM0-HT0179-051099-064-g10&lt3=1999-10-05&lt4=1)  
 Seq primer: puc 18 forward  
 High quality sequence stop: 2.  
 Location/Qualifiers  
 1..550  
 /organism="Homo sapiens"  
 /mol\_type="mRNA"  
 /db\_xref="taxon:9606"  
 /dev\_stage="Adult"  
 /clone\_lib="HT0179"  
 /note="Organ: head neck; Vector: puc18; Site 1: SmaI; Site 2: SmaI; A mini-library was made by cloning products derived from ORSTES PCR (U.S. Letters Patent application No. 196,716 - Ludwig Institute for Cancer Research) profiles into the pUC 18 vector. Reverse transcription of tissue mRNA and cDNA amplification were performed under low stringency conditions."  
 BASE COUNT 179 a 111 c 94 g 166 t  
 ORIGIN  
 Query Match 80.0%; Score 16; DB 13; Length 550;  
 Best local Similarity 100.0%; Pred. No. 81;  
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 Oy 2 CTTAGGAGGACAGT 17  
 Db 279 CTTAGGAGGACAGT 294  
 RESULT 22  
 W52429 554 bp mRNA linear EST 31-MAY-1996  
 LOCUS W52429  
 DEFINITION zc94e08.t1 Pancreatic Islet Homo sapiens cDNA clone IMAGE:338822  
 5', mRNA sequence.  
 ACCESSION W52429  
 VERSION W52429.1 GI:1349780  
 KEYWORDS EST.  
 SOURCE Homo sapiens (human)  
 ORGANISM Homo sapiens  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE 1 (bases 1 to 554)  
 AUTHORS Hillier,L., Lennon,G., Becker,M., Bonaldo,M.F., Chiapelli,B., Chisone,S., Dietrich,N., Dubuque,T., Favello,A., Gish,W., Hawkins,M., Hulman,M., Kucaba,T., Lacy,M., Le,M., Le,N., Maritz,E., Moore,B., Morris,M., Parsons,J., Prange,C., Rikkin,L., Rolling,T., Schellenberg,K., Soares,M.B., Tan,F., Thierry-Mieg,J., Treviskis,E., Underwood,K., Wohlmann,P., Waterson,R., Wilson,R. and Marra,M.  
 Generation and analysis of 280,000 human expressed sequence tags  
 Genome Res. 6 (9), 807-828 (1996)  
 97044478  
 MEDLINE 8889349  
 PUBMED  
 COMMENT Contact: Wilson RK  
 Washington University School of Medicine  
 444 Forest Park Parkway, Box 8501, St. Louis, MO 63108  
 Tel: 314 286 1800  
 Fax: 314 286 1810  
 Email: est@watson.wustl.edu  
 This clone is available royalty-free through LLNL; contact the IMAGE Consortium (info@image.llnl.gov) for further information.  
 Seq primer: mob.REGA+ET  
 High quality sequence stop: 440.  
 Location/Qualifiers  
 1..554  
 /organism="Homo sapiens"  
 /mol\_type="mRNA"  
 /db\_xref="GDB:1264196"  
 /db\_xref="taxon:9606"  
 /clone="IMAGE:338822"  
 /issue\_type="pancreatic islet"  
 /lab\_host="SOLR cells (kanamycin resistant)"  
 /clone\_lib="Pancreatic Islet"  
 /note="Organ: pancreas; Vector: pBluescript SK-; Site 1: EcoRI; Site 2: XhoI; Reference: Hum Mol Gen 2, 1795 (1993) Takeda et al. Cloned unidirectionally. Primer: Oligo dT.  
 -5' adaptor sequence: 5' GAATTCGGACAG 3' -3' adaptor sequence: 5' CTCGAGTTTCTTTTCTTTT 3'"  
 BASE COUNT 181 a 97 c 95 g 180 t 1 others  
 ORIGIN  
 Query Match 80.0%; Score 16; DB 14; Length 554;  
 Best local Similarity 100.0%; Pred. No. 81;  
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 Oy 2 CTTAGGAGGACAGT 17  
 Db 412 CTTAGGAGGACAGT 427  
 RESULT 23  
 B1751017 591 bp mRNA linear EST 25-SEP-2001  
 LOCUS B1751017  
 DEFINITION Ta01\_05d02 C  
 Ta01 AAFc EECORC Fusarium graminearum inoculated wheat heads  
 Triticum aestivum cDNA clone Ta01\_05d02, mRNA sequence.  
 B1751017  
 VERSION B1751017.1 GI:15772819  
 KEYWORDS EST.  
 SOURCE Triticum aestivum (bread wheat)  
 ORGANISM Triticum aestivum  
 Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Pooidae  
 ; Triticeae; Triticum.  
 1 (bases 1 to 591)  
 Ouellet,T., Dan,H., Koul,A., Chapados,T., Couroux,P., De Moors,A., Harris,L.J., Hattori,J.I., Robert,L.S., Singh,J.A., Sprott,D. and Tinker,N.A.  
 Expressed Sequence Tags from Wheat Heads 24 Hours after Spray Inoculation with Fusarium graminearum  
 Unpublished  
 Contact: Ouellet, Therese  
 Eastern Cereal and Oilseed Research Centre  
 Agriculture and Agri-Food Canada  
 Neatby Bldg., Central Experimental Farm, Ottawa, Ontario, KIA 0C6,

CANADA  
Tel: (613) 759-1658  
Fax: (613) 759-1701  
Email: onelett@em.agr.ca.

## FEATURES

source

Location/Qualifiers

1..591

/organism="Triticum aestivum"

/mol\_type="mRNA"

/cultivar="Frontana"

/db\_xref="taxon:4565"

/clone="TA01.05d02"

/tissue\_type="heads"

/dev\_stage="anthesis"

/clone\_lib="TA01\_AAFc\_ECORC\_Fusarium\_graminearum\_inoculate"

d wheat heads"

/note="Vector: pGEM-T easy; Site 1: EcoRI; Site 2: EcoRI; Controlled chamber-grown wheat heads were spray inoculated at mid-anthesis with a Fusarium graminearum macroconidial suspension (50,000 spores/ml) and kept under intermittent misting for 24 hours, then collected and immediately frozen in liquid nitrogen."

162 a 157 c 132 g 140 t

BASE COUNT

ORIGIN

Query Match

Best Local Similarity 80.0%; Score 16; DB 12; Length 591;

Matches 16; Conservativity 100.0%; Pred. No. 82;

Matches 16; Conservativity 0; Mismatches 0; Indels 0; Gaps 0;

Oy

3 TTAGGAGGAGCAAGT 18

Db

272 TTAGGAGGAGCAAGT 257

RESULT 24

HSM095392 standard; RNA; EST; 599 BP.

ID

HSM095392

XX

BX501675;

XX

BX501675.1

SV

XX

09-MAY-2003 (Rel. 75, Created)

DT

09-MAY-2003 (Rel. 75, Last updated, Version 1)

XX

XX

DE

Homo sapiens mRNA; EST DKFZp779F1664\_r1 (from clone DKFZp779F1664)

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EST; expressed sequence tag.

XX

Homo sapiens (human)

OS

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia;

OC

Eutheria; Primates; Catarrhini; Homiidae; Homo.

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/db\_xref="taxon:9606"  
/mol\_type="mRNA"  
/organism="Homo sapiens"  
/clone="DKFZp779F1664"  
/clone\_lib="779 (synonym: hnccl). Vector pSport1\_Sfi; host DH10B; sites SfiI + SfiII"  
/dev\_stage="fetal"  
/tissue\_type="liver"

Sequence 599 BP; 182 A; 105 C; 100 G; 212 T; 0 other;

SQ

Query Match

Best Local Similarity 80.0%; Score 16; DB 2; Length 599;

Matches 16; Conservativity 100.0%; Pred. No. 82; Mismatches 0; Indels 0; Gaps 0;

Oy

2 CTTAGGAGGAGCAAGT 17

Db

479 CTTAGGAGGAGCAAGT 494

RESULT 25

AUI37185

LOCUS

DEFINITION

AUI37185 PLAC1 Homo sapiens CDNA clone PLAC1005960 5', mRNA

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.

REFERENCE

AUTHORS

Ota, T., Nishikawa, T., Suzuki, Y., Ishii, S., Saito, K., Kawai, Y., Yamamoto, J., Wakamatsu, A., Nakamura, Y., Nagai, T., Sugano, S. and

Isogai, T.

HRI human CDNA project

Unpublished

CONTACT: Takao Isogai

Genomics Laboratory

Helix Research Institute

1532-3 Yana, Kisarazu, Chiba 292-0812, Japan

Tel: 81-438-52-3975

Fax: 81-438-52-3986

Email: genom@hri.co.jp

HRI human CDNA project; 5'- &amp; 3'-end one pass sequencing; Helix

Research Institute; CDNA library construction; Department of

Virology, Institute of Medical Science, University of Tokyo, and

Helix Research Institute.

Location/Qualifiers

1..615

/organism="Homo sapiens"

/mol\_type="mRNA"

/db\_xref="taxon:9606"

/clone="PLAC1005960"

/tissue\_type="placenta"

/clone\_lib="PLAC1"

/note="Vector: pME18SFL3"

BASE COUNT

200 a 124 c 92 g 196 t 3 others

ORIGIN

Query Match

Best Local Similarity 80.0%; Score 16; DB 9; Length 615;

Matches 16; Conservativity 100.0%; Pred. No. 83; Mismatches 0; Indels 0; Gaps 0;

Oy

2 CTTAGGAGGAGCAAGT 17

Db

87 CTTAGGAGGAGCAAGT 102

RESULT 26

B1156614

LOCUS

B1156614 630 bp mRNA linear EST 05-JUL-2001

DEFINITION 602921206F1 NCI\_CGAP\_Mam3 Mus musculus cDNA clone IMAGE:5061625 5', mRNA sequence.

ACCESSION B156614

VERSION B156614.1 GI:14616615

KEYWORDS EST.

SOURCE Mus musculus (house mouse)

ORGANISM Mus musculus

REFERENCE NIH-MGC <http://mgc.nci.nih.gov/>.

AUTHORS 1 (bases 1 to 630)

TITLE National Institutes of Health, Mammalian Gene Collection (MGC)

JOURNAL Unpublished

COMMENT Contact: Robert Strausberg, Ph.D.  
Email: [cgapbs-remail.nih.gov](mailto:cgapbs-remail.nih.gov)  
Tissue Procurement: Lothar Hennighausen Ph.D., Chu-Xia Deng Ph.D.  
CDNA Library Preparation: Life Technologies, Inc.  
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNLN)  
DNA Sequencing by: Incyte Genomics, Inc.  
Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LNLN at: <http://image.llnl.gov>  
Plate: LNA11166 row: k column: 02  
High quality sequence stop: 624.  
Location/Qualifiers

FEATURES

source 1..630

/organism="Mus musculus"

/mol\_type="mRNA"

/strain="129, C57BL/6J, FVB/N"

/db\_xref="taxon:10090"

/clone="IMAGE:5061625"

/tissue\_type="tumor, gross tissue"

/dev\_stage="10 months"

/lab\_host="DH10B"

/clone\_lib="NCI\_CGAP\_Mam3"

/note="Organ: mammary; Vector: PCMV-SPORT6; Site 1: SalI; Site 2: NotI; Cloned unidirectionally. Primer: Oligo dt. Library constructed by Life Technologies. Investigators providing samples: Lothar Hennighausen/Chu-Xia Deng, NIH Reference for transgenic model: Xu et al., Nature Genetics 22, 37-43 (1999)."

BASE COUNT 145 a 133 c 166 g 186 t

ORIGIN

Query Match 80.0%; Score 16; DB 12; Length 630;  
Best Local Similarity 100.0%; Pred. No. 83;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 3 TTAGGAGGACACAGTC 18  
|||||  
Db 558 TTAGGAGGACACAGTC 573

RESULT 27  
AV821593 661 bp mRNA linear EST 01-APR-2002  
LOCUS AV821593 RABF4 Arabidopsis thaliana cDNA clone RABF04-13-P03 5', mRNA sequence.  
ACCESSION AV821593  
VERSION AV821593.1 GI:19863621  
KEYWORDS EST.  
SOURCE Arabidopsis thaliana (thale cress)  
ORGANISM Arabidopsis thaliana  
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsids.  
1 (bases 1 to 661)  
Seki, M., Narusaka, M., Ishida, Y., Kamiya, A., Saitou, M., Nakajima, M., Oono, Y., Sakurai, T., Carninci, P., Kawai, J., Itoh, M., Ishii, Y., Arakawa, T., Shibata, K., Shinagawa, A., Muramatsu, M., Hayashizaki, Y. and Shinozaki, K.  
TITLE Large scale analysis of Arabidopsis full-length cDNA (2002b)  
JOURNAL Unpublished

COMMENT Contact: Motoaki Seki  
Plant Functional Genomics Research Group  
RIKEN Genomic Sciences Center  
3-1-1 Koyadai, Tsukuba, Ibaraki 305-0074, Japan  
Tel: 81-298-36-4359  
Fax: 81-298-36-9060  
Email: [mseki@rc.riken.go.jp](mailto:mseki@rc.riken.go.jp)  
An Arabidopsis full-length cDNA library was constructed essentially as reported previously (Seki et al., 1998). This clone is in a modified pBluescript vector as a SrfI/XhoI insert. Please visit our web site ([http://www.gsc.riken.go.jp/e/plant/index\\_e.html](http://www.gsc.riken.go.jp/e/plant/index_e.html)) for further details.

FEATURES

source 1..661

/organism="Arabidopsis thaliana"

/mol\_type="mRNA"

/db\_xref="taxon:3702"

/clone="RABF04-13-P03"

/dev\_stage="rossette plants"

/lab\_host="SOLR"

/clone\_lib="RABF4"

/note="Site 1: SrfI; Site 2: XhoI; subjected to cold-treated(1,2,5,10,24 hr)"

BASE COUNT 188 a 132 c 140 g 201 t

ORIGIN

Query Match 80.0%; Score 16; DB 9; Length 661;  
Best Local Similarity 100.0%; Pred. No. 84;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 2 CTTAGGAGGACACAGT 17  
|||||  
Db 139 CTTAGGAGGACACAGT 154

RESULT 28  
B0273261/c 668 bp mRNA linear EST 09-APR-2002  
LOCUS B0273261 Y. Ogihara unpublished cDNA library, Wh\_on Triticum  
DEFINITION aestivum cDNA clone wholieg03 3', mRNA sequence.  
ACCESSION B0273261  
VERSION B0273261.1 GI:20098087  
KEYWORDS EST.  
SOURCE Triticum aestivum (bread wheat)  
ORGANISM Triticum aestivum  
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Pooidae  
1 (bases 1 to 668)  
Ogihara, Y. and Murai, K.  
TITLE Expressed genes in Triticum aestivum  
JOURNAL Unpublished  
COMMENT Contact: Tadao Shin-i  
Center For Genetic Resource Information  
National Institute of Genetics  
1111 Yata, Mishima, Shizuoka 411-8540, Japan  
Tel: 81-559-81-6856  
Fax: 81-559-81-6855  
Email: [tschini@genes.nig.ac.jp](mailto:tschini@genes.nig.ac.jp).  
Location/Qualifiers

FEATURES

source 1..668

/organism="Triticum aestivum"

/mol\_type="mRNA"

/cultivar="Chinese Spring"

/db\_xref="taxon:4565"

/clone="wholieg03"

/tissue\_type="pistil at heading date"

/dev\_stage="Peekes" scale 10.5"

/clone\_lib="Y. Ogihara unpublished cDNA library, Wh\_on"

BASE COUNT 172 a 185 c 148 g 163 t

ORIGIN

Query Match 80.0%; Score 16; DB 12; Length 668;

Best Local Similarity 100.0%; Pred. No. 84;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 TTAGAGGAGAACAGTC 18  
|||||  
Db 276 TTAGAGGAGAACAGTC 261

RESULT 29  
B0539794

LOCUS B0539794 703 bp mRNA linear EST 27-MAY-2003  
DEFINITION PTM0102 Phaeodactylum tricornutum Uni-Zap XR Phaeodactylum tricornutum cDNA 5', mRNA sequence.

ACCESSION B0539794  
VERSION B0539794.1 GI:21395364  
KEYWORDS EST.

SOURCE Phaeodactylum tricornutum

ORGANISM Phaeodactylum tricornutum  
Eukaryota; stramenopiles; Bacillariophyta; Bacillariophyceae; Bacillariophycidae; Naviculales; Phaeodactylaceae; Phaeodactylum.

REFERENCE 1 (bases 1 to 703)  
Scala, S., Carels, N., Falciatore, A., Chiusano, M.L. and Bowler, C.  
Genome properties of the diatom Phaeodactylum tricornutum  
Plant Physiol. 129 (3), 993-1002 (2002)

AUTHORS  
TITLE  
JOURNAL  
MEDLINE  
PUBMED  
COMMENT

Contact: Bowler C  
Laboratory of Molecular Plant Biology  
Stazione Zoologica 'Anton Dohrn'  
Villa Comunale, I-80121, Napoli, Italy  
Tel: 39 081 583 3268/3211  
Fax: 39 081 764 1355  
Email: chris@alpha.szn.it  
Seq primer: 73 backward

FEATURES  
Location/Qualifiers

1..703

/organism="Phaeodactylum tricornutum"

/mol\_type="mRNA"

/db\_xref="taxon:2850"

/cell\_line="CCMP632"

/clone\_lib="Phaeodactylum tricornutum Uni-Zap XR"

/note="Vector: Uni-Zap XR vector; Site\_1: Eco RI; Site\_2: Xho I"

BASE COUNT 183 a 210 c 182 g 128 t

ORIGIN

Query Match 80.0%; Score 16; DB 13; Length 703;

Best Local Similarity 100.0%; Pred. No. 85;

Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 AGGAGAACAGTCCC 20  
|||||  
Db 359 AGGAGAACAGTCCC 374

RESULT 30  
B0635192

LOCUS B0635192 740 bp mRNA linear EST 23-SEP-2002  
DEFINITION 003D06 Infected Arabidopsis Leaf Arabidopsis thaliana cDNA, mRNA sequence.

ACCESSION B0635192  
VERSION B0635192.1 GI:23302447  
KEYWORDS EST.

SOURCE Arabidopsis thaliana (thale cress)

ORGANISM Arabidopsis thaliana  
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids

REFERENCE 1 (bases 1 to 740)  
Lundsgaard, M., Emmersen, J., Nielsen, K.L., Wilson, I., Somerville, S. and Wellinder, K.G.  
EST sequencing of Erysiphe cichoracearum infected Arabidopsis plants

JOURNAL  
COMMENT

Unpublished  
Contact: Karen G. Wellinder  
Institut for Biotechnologi  
Aalborg Universitet  
Sohnsgaardholmsvej 49, 9000 Aalborg, Denmark  
Tel: +45 96358467  
Fax: +45 98141808  
Email: kgw@bio.auc.dk.

FEATURES  
Location/Qualifiers

1..740

/organism="Arabidopsis thaliana"

/mol\_type="mRNA"

/strain="Columbia"

/db\_xref="taxon:3702"

/dev\_stage="Plant 3 weeks old, three days post infection"

/clone\_lib="Infected Arabidopsis leaf"

/note="Organ: leaf; Vector: plasmid; Mixed cDNA library of Arabidopsis and E. cichoracearum infected leaf from three weeks old Arabidopsis plants. Plants were harvested 3 days after infection and mRNA oligo dt selected."

BASE COUNT 205 a 142 c 170 g 223 t

ORIGIN

Query Match 80.0%; Score 16; DB 13; Length 740;

Best Local Similarity 100.0%; Pred. No. 86;

Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 CTTAGAGAACAGT 17  
|||||  
Db 130 CTTAGAGAACAGT 145

RESULT 31  
B115344

LOCUS B115344 745 bp mRNA linear EST 26-JUN-2001  
DEFINITION 602863159F1 NIH\_MGC\_17 Homo sapiens cDNA clone IMAGE:5022326 5', mRNA sequence.

ACCESSION B115344  
VERSION B115344.1 GI:14566245  
KEYWORDS EST.

SOURCE Homo sapiens (human)

ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.

REFERENCE 1 (bases 1 to 745)  
NIH-MGC <http://mgc.nci.nih.gov/>.

AUTHORS National Institutes of Health, Mammalian Gene Collection (MGC)

TITLE Unpublished

JOURNAL Contact: Robert Strausberg, Ph.D.

COMMENT Email: cgabbe-remail.nih.gov

Tissue Procurement: ATCC

cDNA Library Preparation: Ling Hong/Rubin Laboratory

cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNL)

DNA Sequencing by: Incyte Genomics, Inc.

Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LNL at:

<http://image.llnl.gov>

Plate: LUCM1842 row: e column: 15

High quality sequence stop: 674.

FEATURES  
Location/Qualifiers

1..745

/organism="Homo sapiens"

/mol\_type="mRNA"

/db\_xref="taxon:9606"

/clone="IMAGE:5022326"

/tissue\_type="rhabdomyosarcoma"

/lab\_host="DH10B (phage-resistant)"

/clone\_lib="NIH MGC 17"

/note="Organ: muscle; Vector: pORF7; Site\_1: EcoRI; Site\_2: XhoI; cDNA made by oligo-dT priming.

directionally cloned into EcoRI/XhoI sites using the following 5' adaptor: GGCAAGAG(G). Size-selected >500bp

for average insert size 1.8kb. Library constructed by  
ling Hong in the laboratory of Gerald M. Rubin (University  
of California, Berkeley) using ZAP-cDNA synthesis kit  
(Stratagene) and Superscript II RT (Life Technologies). "

BASE COUNT 151 a 244 c 204 g 146 t

Query Match 80.0%; Score 16; DB 12; Length 745;  
Best Local Similarity 100.0%; Pred. No. 86;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 CTTAGAGGAGAACAGT 17  
|||||  
Db 689 CTTAGAGGAGAACAGT 704

RESULT 32 BE869673 871 bp mRNA linear EST 20-OCT-2000  
LOCUS 601445775F1.NIH\_MGC\_65 Homo sapiens cDNA clone IMAGE:3849618 5',  
DEFINITION mRNA sequence.

ACCESSION BE869673.1 GI:10318358  
VERSION EST.  
KEYWORDS Homo sapiens (human)

SOURCE Homo sapiens  
ORGANISM Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE 1 (bases 1 to 871)  
AUTHORS NIH-MGC http://mgc.nci.nih.gov/.

TITLE Unpublished  
JOURNAL National Institutes of Health, Mammalian Gene Collection (MGC)

COMMENT Contact: Robert Strausberg, Ph.D.  
Email: cgapbs-rc@mail.nih.gov  
Tissue Procurement: ATCC

CDNA Library Preparation: Life Technologies, Inc.  
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNLN)  
DNA Sequencing by: Incyte Genomics, Inc.  
Clone distribution: MGC clone distribution information can be  
found through the I.M.A.G.E. Consortium/LNLN at:  
http://image.llnl.gov

Plate: LLM9567 row: f column: 19  
http://image.llnl.gov

## FEATURES

High quality sequence stop: 676.  
Location/Qualifiers

1..871  
/organism="Homo sapiens"

/mol\_type="mRNA"

/db\_xref="taxon:9606"

/clone="IMAGE:3849618"

/tissue\_type="adenocarcinoma"

/lab\_host="DH10B (phage-resistant)"

/clone\_lib="NIH\_MGC\_65"

/note="Organ: colon; Vector: pCMV-SPORT6; Site 1: NotI;  
Site 2: SalI; Cloned unidirectionally. Primer: Oligo dt.  
Average insert size 1.8 kb. Library constructed by Life  
Technologies."

BASE COUNT 274 a . 169 c 161 g 267 t

Query Match 80.0%; Score 16; DB 10; Length 871;  
Best Local Similarity 100.0%; Pred. No. 89;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 CTTAGAGGAGAACAGT 17  
|||||  
Db 180 CTTAGAGGAGAACAGT 195

RESULT 33

LOCUS B1088175 872 bp mRNA linear EST 20-JUN-2001  
DEFINITION 602851213F1.NIH\_MGC\_10 Homo sapiens cDNA clone IMAGE:4992843 5',  
mRNA sequence.

ACCESSION B1088175  
VERSION B1088175.1 GI:14506505  
KEYWORDS EST.  
SOURCE Homo sapiens (human)

ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE 1 (bases 1 to 872)  
AUTHORS NIH-MGC http://mgc.nci.nih.gov/.

TITLE Unpublished  
JOURNAL National Institutes of Health, Mammalian Gene Collection (MGC)

COMMENT Contact: Robert Strausberg, Ph.D.  
Email: cgapbs-rc@mail.nih.gov  
Tissue Procurement: ATCC  
CDNA Library Preparation: Life Technologies, Inc.  
CDNA Library Arrayed by: Incyte Genomics, Inc.  
DNA Sequencing by: Incyte Genomics, Inc.  
Clone distribution: MGC clone distribution information can be  
found through the I.M.A.G.E. Consortium/LNLN at:  
http://image.llnl.gov

Plate: LLM11012 row: i column: 04  
http://image.llnl.gov

High quality sequence stop: 738.  
Location/Qualifiers

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/organism="Homo sapiens"

/mol\_type="mRNA"

/db\_xref="taxon:9606"

/clone="IMAGE:4992843"

/cell\_line="MGC36"

/lab\_host="DH10B"

/clone\_lib="NIH\_MGC\_10"

/note="Organ: cervix; Vector: pCMV-SPORT6; Site 1: NotI;  
Site 2: SalI; Cloned unidirectionally. Primer: Oligo dt.  
Average insert size 1.5 kb. Library prepared by Life  
Technologies."

BASE COUNT 281 a 165 c 151 g 275 t

Query Match 80.0%; Score 16; DB 12; Length 872;  
Best Local Similarity 100.0%; Pred. No. 89;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 CTTAGAGGAGAACAGT 17  
|||||  
Db 130 CTTAGAGGAGAACAGT 145

RESULT 34

LOCUS BUI77008 879 bp mRNA linear EST 04-SEP-2002  
DEFINITION AGENCOURT\_7940818.NIH\_MGC\_71 Homo sapiens cDNA clone IMAGE:6155256  
5', mRNA sequence.

ACCESSION BUI77008  
VERSION BUI77008.1 GI:22690992  
KEYWORDS EST.  
SOURCE Homo sapiens (human)

ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE 1 (bases 1 to 879)  
AUTHORS NIH-MGC http://mgc.nci.nih.gov/.

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CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNLN)  
DNA Sequencing by: Agencourt Bioscience Corporation  
Clone distribution: MGC clone distribution information can be  
found through the I.M.A.G.E. Consortium/LNLN at:  
http://image.llnl.gov

Plate: LLM13497 row: k column: 01

FEATURES High quality sequence stop: 691.  
Location/Qualifiers

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/db\_xref="taxon:9606"  
/clone="IMAGE:6155256"  
/tissue\_type="leiomyosarcoma"  
/lab\_host="DH10B (phage-resistant)"  
/cloning\_lib="NIH\_MGC\_71"  
/note="Organ: uterus; Vector: pCMV-Sport6; Site\_1: NotI;  
Site\_2: SalI; Cloned unidirectionally. Primer: Oligo dT.  
Average insert size 2.1 kb."  
BASE COUNT 280 a 159 c 143 g 297 t  
ORIGIN

Query Match 80.0%; Score 16; DB 13; Length 879;  
Best Local Similarity 100.0%; Pred. No. 89;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 2 CTTAGAGGAACT 17  
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Db 439 CTTAGAGGAACT 454

RESULT 35  
BG740602 884 bp mRNA linear EST 15-MAY-2001  
LOCUS 602631028F1 NCI\_CGAP\_Skn3 Homo sapiens cDNA clone IMAGE:4776247 5',  
DEFINITION mRNA sequence.  
ACCESSION BG740602  
VERSION BG740602.1 GI:14051255  
KEYWORDS EST.  
SOURCE Homo sapiens (human)  
ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.  
1 (bases 1 to 884)  
NIH-MGC http://mgs.nci.nih.gov/  
National Institutes of Health, Mammalian Gene Collection (MGC)  
Unpublished  
Contact: Robert Strausberg, Ph.D.  
Email: cgapbs-r@mail.nih.gov  
Tissue Procurement: James Cleaver, M.D.  
cDNA Library Preparation: Life Technologies, Inc.  
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNL) DNA  
Sequencing by: Incyte Genomics, Inc.  
Clone distribution: MGC clone distribution information can be  
found through the I.M.A.G.E. Consortium/LNL at:  
http://image.llnl.gov  
plate: LLM10628 row: h column: 08  
High quality sequence stop: 753.  
Location/Qualifiers

FEATURES  
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/clone="IMAGE:4776247"  
/lab\_host="DH10B (T1 phage-resistant)"  
/cloning\_lib="NCI CGAP\_Skn3"  
/note="Organ: skin; Vector: pCMV-Sport6; Site\_1: NotI;  
Site\_2: SalI; Cloned unidirectionally. Primer: Oligo dT.  
Average insert size 1.5kb. Library constructed by Life  
Technologies. Note: this is a NCI\_CGAP Library."  
BASE COUNT 264 a 180 c 154 g 286 t  
ORIGIN

Db 193 CTTAGAGGAACT 208

Search completed: August 15, 2003, 10:57:48  
Job time: 1261 secs

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Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
OY 2 CTTAGAGGAACT 17  
|||||